

Transmitted Via Electronic Mail

June 2₄, 2016

Ms. Alice Yeh Remedial Project Manager U.S. Environmental Protection Agency, Region II Emergency and Remedial Response Division 290 Broadway, 19th Floor New York, NY 10007-1866

Re: Combined Sewer Overflow/Stormwater Outfall Program Phase I Evaluation and Recommendation Report Revision 2

Dear Ms. Yeh:

Please find enclosed CSO/SWO Phase I Evaluation and Recommendation Report, Revision 2 dated June 2016 and prepared in accordance with the Combined Sewer Overflow (CSO) / Stormwater Outfall (SWO) Quality Assurance Project Plan (QAPP) Revision 2 of the Recommendation Report incorporates changes resulting from comments contained in the following documents:

- Phase I Data Usability Report Aug. 2014 EPA Comments USEPA provided initial comments on the DQUAR on August 6th, 2015.
- Tierra Responses to USEPA Comments on DQUAR_Final 09.18.15 Tierra submitted a Response to Comment (RTC) document on September 18, 2015, which provided responses to USEPA comments received on August 6th, 2015.
- CDM Smith Comments_CSO -SWO Phase I Tierra Response to Comments CDM Smith (EPA Contractor) submitted final comments to the DQUAR on November 12, 2015.
- Revised Report Review 05.06.2016- CDM Smith submitted comments on Revision 1 of the Recommendation Report submitted April 1, 2016.

If you have any questions regarding the attached report, please feel free to contact me at 732-246-5920.

Sincerely,

Brian Mikucki

Bi think h.

Senior Environmental Scientist

On behalf of Occidental Chemical Corporation

(as successor to Diamond Shamrock Chemicals Company)

Enclosures

Cc: Enrique Castro, Tierra Solutions, Inc.
Clifford Firstenberg, Tierra Solutions, Inc.
Diane Waldschmidt, Environmental Data Services
Kavin Gandhi, ARCADIS
Alain Hebert, ARCADIS

Combined Sewer Overflow/Stormwater Outfall Investigation

Phase I Evaluation/ Recommendation Report

Tierra Solutions, Inc.

East Brunswick, New Jersey

June 2016

Revision 2

Acre	onyms	and Abbreviations	V	
1.	Intro	oduction	1-1	
	1.1	Organization of Report	1-2	
2.	Sum	nmary of Field Activities	2-1	
	2.1	2-1		
	2.2	Mobilization for Sample Collection	2-3	
	2.3	2.3 Sample Collection – Clay Street Combined Sewer Overflow		
	2.4	Decontamination/Cleaning	2-5	
3.	Sum	nmary of Evaluation Process	3-1	
4.	lmpl	lementation Evaluation	4-1	
	4.1	Implementation Requirements and Challenges	4-1	
	4.2	Evaluation of Sampling Methods	4-3	
		4.2.1 High-Solids Mass	4-3	
		4.2.1.1 High-Solids Mass Particulate	4-3	
		4.2.1.2 High-Solids Mass Dissolved	4-5	
		4.2.2 Low-Solids Mass	4-6	
		4.2.2.1 Low-Solids Mass Bulk Sample Collection	4-6	
		4.2.2.2 Low-Solids Mass Bulk Laboratory Filtration	4-7	
		4.2.3 Whole Water	4-10	
		4.2.4 Grab Metals	4-11	
	4.3	Summary of Implementability Evaluation	4-12	
5.	Ana	lytical Data Evaluation	5-1	
	5.1	Data Usability	5-1	
	5.2	Decontamination	5-2	
	5.3	5.3 Field Blank Results and Affected Sample Results		
	5.4	Steps 3 and 4: Frequency of Detections	5-3	
		5.4.1 Polychlorinated Dibenzo-p-dioxins/Polychlorinated Dibenzofurans	5-3	

		5.4.2	Polychlorinated Biphenyl Congeners	5-4
		5.4.3	Aroclor Polychlorinated Biphenyls	5-5
		5.4.4	Organochlorine Pesticides	5-5
		5.4.5	Semivolatile Organic Compounds	5-6
		5.4.6	Semivolatile Organic Compounds Select Ion Monitoring	5-7
		5.4.7	Chlorinated Herbicides	5-8
		5.4.8	Cyanide	5-9
		5.4.9	Volatile Organic Compounds	5-9
		5.4.10	Total Extractable Petroleum Hydrocarbons	5-10
	5.5	Impacts	s of Achieved Analytical Sensitivity	5-10
	5.6	Addition	nal Data Evaluation	5-11
6.	Con	clusion/	Recommendation	6-1
7.	Refe	rences		7-1
Tab	oles			
	2-1	Sur	mmary of Samples Collected and Analyzed	2-4
	3-1	Ana	alytical Groups Included in Phase I Evaluation Process	3-1
	4-1	LSI	M Bulk Liquid Volume Requirements by Analtyical Group	4-8
	4-2		geted LSM Dissolved Volume and Corresponding Actual LSM Bulk Volume Filtered by alytical Group	4-9
	4-3	Tar Gro	rgeted LSM Particulate Mass and Corresponding Actual LSM Particulate Mass by Analytic oup	cal 4-9
	5-1	Sur	nmary of Data Quality Failures	5-1
	5-2	Red	commended Sample Collection Method – PCDDs/PCDFs	5-4
	5-3	Red	commended Sample Collection Method – PCB Congeners	5-5
	5-4	Red	commended Sample Collection Method – Aroclor PCBs	5-5
	5-5	Red	commended Sample Collection Method – Organochlorine Pesticides	5-6
	5-6	Red	commended Sample Collection Method – SVOCs	5-7
	5-7	Red	commended Sample Collection Method – SVOCs SIM	5-7

5-8	Recommended Sample Collection Method – Chlorinated Herbicides	5-9
5-9	Impact of PQL Exceedances	5-11
6-1	Phase I Sample Collection Method Recommendations	6-1
Figures		
2-1	CSO/SWO Sample Collection System and Schematic	
2-2	CSO/SWO Sample Collection System and Schematic – Cross-Section A and B	
2-3	CSO/SWO Sample Collection System and Schematic – Cross-Section C	
2-4	Schematic of Weighted Rod/Tubing Assembly	
3-1	Phase I Evaluation Process Flow Chart (embedded in text)	
Appendices		
Α	Event #1, Attempt #1 Results – PCDDs/PCDFs	
В	Event #1, Attempt #1 Results – PCB Congeners	
С	Contingency Samples Used During CSO Phase I Sampling Events	
D	CSO/SWO Phase I Field Blank Contamination Results	
Е	Field Blank Results Assessment	
F	Detailed Evaluation Sheets (Worksheet #11) - PCDDs/PCDFs	
G	Detailed Evaluation Sheets (Worksheet #11) - PCB Congeners	
Н	Detailed Evaluation Sheets (Worksheet #11) - Aroclor PCBs	
I	Detailed Evaluation Sheets (Worksheet #11) - Organochlorine Pesticides	
J	Detailed Evaluation Sheets (Worksheet #11) - SVOCs	
K	Detailed Evaluation Sheets (Worksheet #11) - SVOCs SIM	
L	Detailed Evaluation Sheets (Worksheet #11) - Chlorinated Herbicides	
М	Detailed Evaluation Sheets (Worksheet #11) - Cyanide	
N	Detailed Evaluation Sheets (Worksheet #11) - VOCs	
0	Detailed Evaluation Sheets (Worksheet #11) - TEPH	
Р	CSO/SWO Phase I Data Quality Usability Assessment Report	

Attachment

1 Phase I Report Addendum – Additional Data Evaluation

Acronyms and Abbreviations

CFC continuous flow centrifuge

CH clean hands

COC constituent of concern

COPC constituent of potential concern

COPEC constituent of potential ecological concern

CSO combined sewer overflow

CSO/SWO S&AP Combined Sewer Overflow/Stormwater Overflow Sampling and

Analytical Plan

DH dirty hands

DOC dissolved organic carbon

EDL estimated detection limit

HSM high-solids mass

LPRSA Lower Passaic River Study Area

LSM low-solids mass

MDL method detection limit

mg/L milligrams per liter

NOAA's NWS National Oceanic and Atmospheric Administration's National Weather

Service

PCB polychlorinated biphenyl

PCDD polychlorinated dibenzo-p-dioxin

PCDF polychlorinated dibenzofuran

Phase I Report Phase I Evaluation/Recommendation Report

POC particulate organic carbon

PQL project quantitation limit

POTW publicly owned treatment works

PVSC Passaic Valley Sewerage Commission

QA quality assurance

CSO/SWO Investigation QAPP Combined Sewer Overflow/Stormwater Outfall Investigation Quality

Assurance Project Plan

QC quality control

SIM selective ion monitoring

SOP standard operating procedure

SVOC semivolatile organic compound

SWO stormwater outfall

TAL Target Analyte List

TDS total dissolved solid

TEPH total extractable petroleum hydrocarbons

Tierra Solutions, Inc.

TOC total organic carbon

TSS total suspended solids

USEPA United States Environmental Protection Agency

VOC volatile organic compound

Revision Data: June 2016

1. Introduction

This Phase I Evaluation/Recommendation Report (Phase I Report) has been developed by Tierra Solutions, Inc. (Tierra), on behalf of Occidental Chemical Corporation, the successor to Diamond Shamrock Chemicals Company (formerly known as Diamond Alkali Company). This Phase I Report documents the evaluation of data collected as part of Phase I of the combined sewer overflow/stormwater outfall (CSO/SWO) investigation implemented under the U.S. Environmental Protection Agency- (USEPA-) approved Combined Sewer Overflow/Stormwater Outfall Investigation Quality Assurance Project Plan (QAPP; Tierra 2013). The QAPP was developed to guide the collection of CSO, SWO, and publicly owned treatment works (POTW) samples from within the Lower Passaic River Study Area (LPRSA). The main objective of the CSO/SWO investigation is to characterize and quantify contaminants in both particulate- and dissolved-phases present in runoff discharging to the LPRSA via CSO and SWO conveyances, such that subsequent determinations of contaminant loadings can be made using models, developed by others, for the lower Passaic River.

The unique challenge of the CSO/SWO investigation is the quantification of organic contaminants found in the effluent of CSOs and SWOs, which are typically bound to particulates and, to a lesser degree, in the dissolved-phase. Quantitation limits associated with the particulate-phase of the effluent are particularly challenging to achieve, in that quantitation limits needed to reach the program data quality objectives require a sufficient mass of solids be collected for detection via standard, USEPA-approved laboratory analyses. The challenges associated with collecting a sufficient mass of solids for analysis are one of the focuses of the Phase I investigation.

Various sampling methods have been used previously in the LPRSA to collect the necessary solids mass for analysis, with varying results. As such, a two-phased approach for the CSO/SWO investigation was developed in coordination with the USEPA. This two-phased approach incorporates, as Phase I, an initial side-by-side sampling program for evaluating three sampling approaches to inform the selection of the most appropriate sampling approach to quantify contaminants in the solid- (particulate), dissolved-, and whole water-phases: low-solids mass (LSM), high-solids mass (HSM), and whole water. Phase II of the program will consist of collecting CSO, SWO, and POTW samples at target locations using the sampling and analytical technique(s) selected after evaluation of Phase I results (the subject of this Phase I Report).

The LSM approach is a modification of the methods described in the USEPA Combined Sewer Overflow/Stormwater Overflow Sampling and Analytical Plan, Revision No. 2.0, August 2008 (CSO/SWO S≈ USEPA 2008). The CSO/SWO S&AP was, in turn, based on methods that were implemented in the 1998 to 2004 Contaminant Assessment and Reduction Program (Great Lakes Environmental Center 2008) and the 2008 USEPA CSO/SWO solid-phase sampling conducted by Malcolm Pirnie, Inc. (2008). The LSM approach requires modifications to standardized analytical methods for solids sample analyses because a relatively small mass of particulates is acquired during the sample collection procedure. The HSM approach was proposed in the LPRSA Remedial Investigation – Combined Sewer Overflow Investigation, Volume 1,

Work Plan/Field Sampling Plan Revision No. 1 (Tierra 2002). The HSM approach calls for the collection of a greater mass of particulates than the LSM method, and similar to the mass specified in standardized analytical methods. The whole water approach is similar to the LSM approach, except that the particulate and dissolved-phases are not separated prior to analysis.

1.1 Organization of Report

The	e remainder of this Phase I Report is organized as follows:
Amenda	Section 2 – Summary of Field Activities: Summarizes the three sample collection methods and associated sample collection activities completed.
desirab	Section 3 – Summary of Evaluation Process: Summarizes the process used to evaluate the implementability and effectiveness of the three sample collection methods.
-montal	Section 4 – Implementation Evaluation: Summarizes the evaluation of the implementability of the three sample collection methods.
residents	Section 5 – Analytical Data Evaluation: Summarizes the evaluation of the analytical data obtained for the three sample collection methods.
randonos	Section 6 – Conclusions/Recommendations: Summarizes the conclusions of the data evaluation process and provides the recommended path forward.
-	Section 7 – References: Provides a summary of the references used in this Phase I Report

Revision Data: June 2016

2. Summary of Field Activities

Phase I sampling consisted of collecting and analyzing samples using three sample collection methods (LSM, HSM, and whole water) during two precipitation events at the selected CSO (Clay Street in Newark, New Jersey). The field sample collection activities were implemented in accordance with the Field Standard Operating Procedures (SOPs) contained in the QAPP (Tierra 2013). It should be noted that the QAPP originally specified collection of samples from two different CSO locations: Clay Street CSO in Newark, New Jersey and Ivy Street CSO in Kearny, New Jersey. However, due to access limitations to the Ivy Street CSO imposed by the City of Kearny and to meet the Phase I implementation schedule, the USEPA and Tierra decided to collect an additional sample at the Clay Street CSO (for a total of two) in lieu of sampling at the Ivy Street CSO during Phase I. Modifications were made to the QAPP (Tierra 2013) to address this change.

2.1 Sample Collection System

A sample collection system was designed to collect all three sample types (LSM, HSM, and whole water) simultaneously from the same effluent stream and over the same period of time by controlling the flow rate of effluent entering different sample collection tanks and the continuous flow centrifuge (CFC). The sample collection system utilized an enclosed trailer as a secure platform for mounting/housing the sampling equipment and controls. Sampling equipment included a bulk sample collection tank, peristaltic pumps (one large-diameter peristaltic pump and three small-diameter perist altic pumps), CFC, and associated tubing and fittings. A stand-alone tow-behind generator was staged near the sample collection trailer during sample collection. Figures 2-1, 2-2, and 2-3 present the schematic of the sample collection equipment setup. SOP No. 2 – Pre-Mobilization and SOP No. 3 – Mobilization, Bulk Sample Collection, and Transportation (Tierra 2013) provide additional details regarding the sample collection system.

During each sampling event, a weighted rod/tubing assembly (Figure 2-4) was deployed into the manhole of the diversion chamber at the Clay Street CSO for bulk sample collection. Large-diameter intake tubing (i.e., 1.125-inch outside diameter for large-diameter high-flow peristaltic pump) was secured to the weighted rod/tubing assembly and connected to a large-diameter high-flow peristaltic pump in the trailer to pump bulk sample for collection. Three sample ports were installed along the large-diameter intake tubing, two before, and one after the CFC. Small-diameter sample tubing and small-diameter peristaltic pumps were connected to the sample ports to pump bulk sample from the large-diameter intake tubing line into two bulk sample collection tanks (whole water/LSM and HSM dissolved bulk sample collection tanks). From an initial single sample flow stream, flow was continuously diverted to the Teflon®-lined (double-lined) whole water/LSM bulk sample collection tank (via the second sample port to generate the LSM and whole water samples) and the CFC (to generate solids in the centrifuge for HSM particulate analysis and CFC effluent for HSM dissolved analysis). A portion of the CFC effluent that passed through the CFC was diverted via the third sample port to the Teflon®-lined (double-lined) HSM dissolved bulk sample collection tank to generate HSM dissolved samples. The flow rate to each bulk sample collection tank was controlled so that the whole water/LSM bulk sample collection tank filled in approximately the same time as the HSM dissolved bulk sample collection

Revision Data: June 2016

tank. The excess effluent that passed through the CFC was returned to the same manhole via largediameter tubing downstream of the CFC and HSM dissolved bulk sample collection tank.

The effluent entered the CFC from the bottom through a stationary feed nozzle and is directed towards the CFC bowl. A variable frequency drive mounted on the trailer was used to operate and control the speed of the CFC. Solids in the bulk effluent were forced to the bowl wall by centrifugal force. The interior of the CFC bowl was lined with a Teflon[®] liner to capture the separated solids. The clarified liquid was continuously discharged through the top of the centrifuge.

Following collection of effluent into the bulk sample collection tanks, aqueous (LSM bulk, HSM dissolved, and whole water) samples were collected using small-diameter pe ristaltic pumps and dedicated Teflon® tubing from the bulk sample collection tanks. The LSM bulk samples were further processed in analytical laboratories, via filtration, to generate LSM particulate and LSM dissolved samples for analysis. HSM particulate samples were collected from the solids retained inthe CFC bowl and liner for laboratory analysis. SOP No. 4 – Sample Processing and Collection (Tierra 2013) provides additional details on sample processing.

Upon receipt of LSM bulk samples by the laboratory, the equipment and procedures described in SOP No. L-24 – LSM Bulk Sample Filtration (Tierra 2013) were utilized to filter the LSM bulk sample, thereby generating LSM particulate and LSM dissolved samples for analysis. Post-filtration of the LSM bulk sample, particulate material captured on the filter media was put forward for analysis as the LSM particulate sample, while the filtrate was analyzed as the corresponding LSM dissolved sample. Two approaches were included in SOP No. L-24 – LSM Bulk Sample Filtration to filter the LSM bulk samples. The primary approach involved the use of pressurized filtration and a flat glass fiber filter(s). The secondary approach utilized a system by which bulk sample is pumped through a wound glass fiber filter cartridge and a flat glass fiber filter in series. The secondary approach was included for use as a contingency when/if excessive clogging was observed during implementation of the primary approach due to sample particulate mass characteristics, such as high total suspended solids (TSS) content or large individual particulate size.

During bulk sample collection at the manhole, TSS/total dissolved solids (TDS) grab samples were collected every 30 minutes via the first sample withdrawal port installed along the large-diameter intake tubing prior to the CFC and whole water/LSM bulk sample collection tank. Additionally during sample collection, selected physiochemical water quality parameters (conductivity, turbidity, and temperature) were measured (logged continuously and manually recorded every 30 minutes using a water quality meter), water depth was measured at the sample collection manhole, and flow data were recorded. An in-line flow meter, located downstream of the CFC, was used to monitor and record flow rate approximately every 30 minutes.

Grab metals samples (including mercury and methyl mercury) were collected in accordance with SOP No. 5 – Metals Sampling via Method 1669 Sampling Ambient Water for Trace Metals at EPA Water Quality Criteria Levels (USEPA 1996) (Tierra 2013). This methodology has been developed based on USEPA Method 1669: Sampling Ambient Water for Trace Metals at EPA Water Quality Criteria Levels (USEPA

1996). Grab (total and dissolved)samples for trace metals analysis, including mercury and methyl mercury, and a TSS sample were collected directly from the manhole into laboratory-supplied containers using a separate peristaltic pump and laboratory-supplied Teflon® tubing. This sampling method was employed so that metals samples could be collected using "clean hands" (CH) and "dirty hands" (DH) sampling methods that minimize potential sample contamination from trace metalsduring sample collection. Sampling activities were conducted with care to minimize exposure of the sample to atmospheric, human, and other sources of potential metals contamination. Dissolved metals samples were collected first by field-filtering (via an in-line filter) the effluent followed by collection of samples for total metals analysis.

2.2 Mobilization for Sample Collection

During Phase I, Tierra conducted weather monitoring on a daily basis using multiple sources to evaluate timing of mobilization for sample collection. For a precipitation event to trigger mobilization for sample collection, the event must have anticipated to produce at least 0.2 inch of rain with an average intensity of at least 0.03 inch per hour with no more than 4 consecutive dry hours during the event. Following a decision to mobilize for sample collection, staff mobilized the sample collection system to the sampling location. Tierra coordinated/communicated with Passaic Valley Sewerage Commission (PVSC) to determine timing of the regulator gate valve closing at the Clay Street CSO and appropriate time for initiating sample collection. Sample collection was only initiated after PVSC confirmed that the regulator gate valve was closed at the Clay Street CSO and that an overflow was occurring. In addition, a sidewalk occupancy permit was obtained in advance from the City of Newark to stage the sample collection system along the sidewalk at the Clay Street CSO; the Newark Police Department were also contacted to provide traffic control. Following bulk sample collection, the sample collection system was transported back to the processing facility at 80 Lister Avenue in Newark, New Jersey. Samples were shipped to analytical laboratories the day after bulk sample collection in accordance with the procedures outlined in the QA PP (Tierra 2013).

2.3 Sample Collection - Clay Street Combined Sewer Overflow

Phase I sampling was completed at the Clay Street CSO between June 2013 and April 2014. It was critical that sufficient sample mass and/or volume be obtained to accomplish the primary objective of this phase: the evaluation and selection of the most appropriate sampling method for each analytical group. For this reason, an analytical hierarchy was established for sample collection. For a given sampling event, if sufficient volume was obtained to complete sampling via the three methods for the analytical groups and matrices, then samples were generated in the sequence described in the analytical hierarchy detailed in the QAPP (Tierra 2013) (with the exception of samples for volatile organic compound [VOC] analysis, which were collected first). In addition to the sample mass/volume required for primary sample analysis (including quality assurance/quality control [QA/QC] samples) contingency sample mass/volume was collected and shipped to the laboratories to mitigate any potential issues related to sample breakage/loss during sample shipment and analysis. Multiple attempts were needed during each sampling event at the Clay Street CSO to collect all samples (primary and contingency) for the target analytical groups using the three sampling approaches.

Table 2-1 summarizes the number and type of samples collected and analyzed during each sampling event/attempt as part of the Phase I sampling program.

Table 2-1
ummary of Samples Collected and Analyzed

Event and	Sample	Date	Collection Method and Analytical Parameters*				
Attempt	Identification		HSM	LSM	Whole Water		
Event #1, Attempt #1 ^b	PR1CSOCLY**-01A PR1**DUP-01A	June 10, 2013	PCDDs/PCDFs, PCB congeners	PCDDs/PCDFs, PCB congeners	PCDDs/PCDFs, PCB congeners, metals, mercury, and methyl mercury		
Event #1, Attempt #2	PR1CSOCLY**-01B PR1**-DUP-01B	July 1, 2013	Alla, excluding PCDDs/PCDFs, PCB congeners, POC, grain size, metals, mercury and methyl mercury	Alla, excluding PCDDs/PCDFs, PCB congeners, TOC, grain size, VOCs, cyanide, TEPH, metals, mercury and methyl mercury	Alla, excluding PCDDs/PCDFs, PCB congeners, DOC, POC, metals, mercury and methyl mercury		
Event #1, Attempt #3°	PR1CSOCLY**-01C PR1**-DUP-01C	April 30, 2014	PCDDs/PCDFs, PCB congeners, chlorinated herbicides	PCDDs/PCDFs, PCB congeners, chlorinated herbicides	PCDDs/PCDFs, PCB congeners, chlorinated herbicides		
Event #2, Attempt #1	PR1CSOCLY**-02A PR1**-DUP-02A	October 7, 2013	VOCs	-	VOCs		
Event #2, Attempt #2 ^b	PR1CSOCLY**-02B PR1**-DUP-02B	December 7, 2013	Alla, excluding VOCs, grain size, POC, metals, mercury and methyl mercury	Alla, excluding VOCs, TOC, grain size, cyanide, TEPH, metals, mercury and methyl mercury	All ^a , excluding VOCs, DOC, POC		

Notes:

- a. All includes the following analyses: polychlorinated dibenzo-p-dioxins/polychlorinated dibenzofurans (PCDDs/PCDFs), polychlorinated biphenyl (PCB) congeners, Aroclor PCBs, organochlorine pesticides, semivolatile organic compounds (SVOCs), SVOC selective ion monitoring (SIM), chlorinated herbicides, metals, mercury, methyl mercury, cyanide, VOCs, total extractable petroleum hydrocarbons (TEPH), TSS, TDS, total organic carbon (TOC), particulate organic carbon (POC), dissolved organic carbon (DOC), and grain size.
- b. Grab total and dissolved metals (including mercury and methyl mercury) samples were collected on June 10, 2013 (Event #1, Attempt #1) and December 7, 2013 (Event #2, Attempt #2).
- c. During Event #1, Attempt #1, two types of solid material ("fine" and "non-fine paper like material") were recovered in the centrifuge bowl. To be consistent with sediment homogenization implemented in subsequent events/attempts (i.e., "fines" and "non-fines" were combined and homogenized), PCDDs/PCDFs and PCB congener samples were collected during Event #1, Attempt #3 (which occurred after both Event #2 attempts) to replace the Event #1, Attempt #1 PCDDs/PCDFs and PCB congener results. In addition, chlorinated herbicides were collected during Event #1, Attempt #3 to obtain an additional set of herbicide data due to a labor atory error identified during the herbicide analysis of the HSM parti culate sample. Laboratory results indicated that a laboratory c ontrol sample associated with the herbicide data had failed during Event #2, Attempt #2.
- * Grab TSS/TDS samples were collected every 30 minutes during each sampling event/attempt in addition to the TSS/TDS samples collected as part of HSM, LSM, and whole water sampling methods.
- ** = Two-character code to indicate sample matrix (e.g., "HP" for HSM particulate).
- = sample not collected/analyzed.

The PCDDs/PCDFs, PCB congeners, and organochlorine pesticides were analyzed by Vista Analytical in El Dorado Hills, California. Brooks Rand laboratory in Seattle, Washington analyzed the total and dissolved metals (including mercury and methyl mercury) samples. The remainder of the analyses was performed by TestAmerica in Burlington, Vermont.

Revision Data: June 2016

2.4 Decontamination/Cleaning

Applicable decontamination procedures were followed throughout the Phase I sample collection program in accordance with SOP No. 6: Decontamination included in the QAPP (Tierra 2013). Between sampling events, a full decontamination of the sample collection system was performed in accordance with Section 2.2.2 of SOP No. 6: Decontamination, included in the QAPP (Tierra 2013). Field sampling equipment designated for analyses other than trace metals (i.e., CFC bowl, CFC bowl Teflon® liner, CFC components, stainless steel fittings, and stainless steel tools used for HSM particulate sample collection) was decontaminated prior to the first sampling attempt for each event. Dedicated sampling equipment (i.e., CFC bowl Teflon® liner, Teflon® tank liners, and small- and large-diameter Teflon® sample tubing) were replaced with new dedicated sampling equipment between events.

Between sampling attempts (e.g., between Attempts #1 and #2 of Event #1), non-dedicated sampling equipment used for HSM particulate sample collection (e.g., CFC bowl, CFC bowl Teflon® liner, CFC components, stainless steel bowls and spoons) was fully decontaminated in accordance with Section 2.2.3 of SOP No. 6, included in the QAPP (Tierra 2013). Note that permanently attached stainless steel fittings associated with the sampling system prior to entry into the CFC bowl were not fully decontaminated; however, a "gross cleaning" procedure was followed as per SOP No. 6 by circulating deionized water through the system. Dedicated sampling equipment (Teflon® tank liners and Teflon® tubing) were not replaced between sampling attempts (unless damaged) as per SOP No. 6.

3. Summary of Evaluation Process

Phase I data was evaluated, on an analytical group basis, for each sampling approach using the following criteria as defined in the QAPP (Tierra 2013):

- Implementability of field sampling and sample processing activities
- Ability to generate sample mass/volume to accommodate the full target analytical groups
- Ability of laboratories to generate usable data
- Ability to generate greater frequency of detection for analytes that are constituents of potential concern (COPCs) and/or constituents of potential ecological concern (COPECs) listed in the Lower Eight Miles of the Lower Passaic River Feasibility Study Report (The Louis Berger Group 2014)
- Ability to generate greater frequency of detection for analytes within a given analytical group.

Analytical groups included in the evaluation were limited to those where samples were collected using two or more of the sampling methods (HSM, LSM, and/or whole water); therefore, the Phase I evaluation process included comparison of the analytical groups as defined in Table 3-1 below.

Table 3-1 Analytical Groups Included in Phase I Evaluation Process

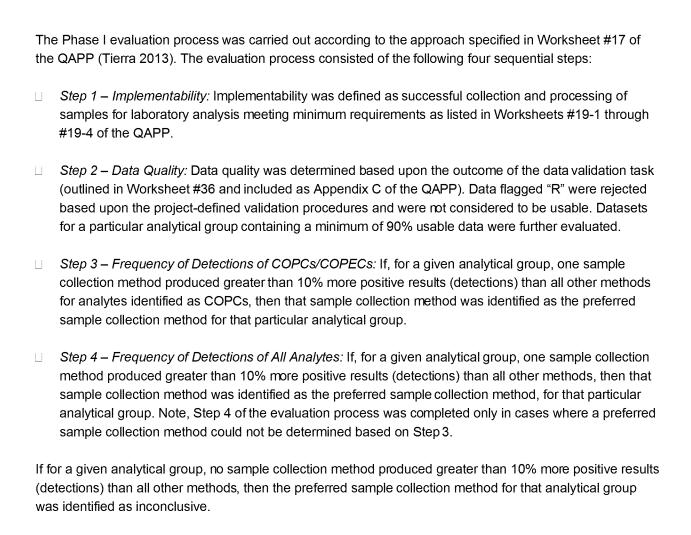
Analytical Groups Included in Phase I Evaluation Process Sampling Methods Implemented Analytical Group								
	Sampli	ng Methods Imp	lemented	Included in Phase I				
Analytical Group	HSM	LSM	Whole Water	Evaluation Process?				
PCDDs/PCDFs	Х	Х	Х	Yes				
PCB Congeners	Х	Х	Х	Yes				
Aroclor PCBs	Х	Х	Х	Yes				
Organochlorine Pesticides	Х	Х	Х	Yes				
SVOCs	Х	Х	Х	Yes				
SVOC SIM	Х	Х	Х	Yes				
Chlorinated Herbicides	Х	Х	Х	Yes				
Cyanide	Х	-	Х	Yes				
VOCs	Х	-	Х	Yes				
TEPH	Х	-	Х	Yes				
TSS	Х	Х	Х	No				
TDS	Х	Х	Х	No				
TOC	Х	-	Х	No				
POC	-	Х	-	No				
DOC	Х	Х	-	No				
Grain Size	-	-	Х	No				
Metals	-	-	Х	No				
Mercury	-	-	Х	No				
Methyl mercury	-	-	Х	No				

Notes:

x = analytical sampling method was performed

^{- =} analytical sampling method was not performed

Revision Data: June 2016



The evaluation process is represented below.

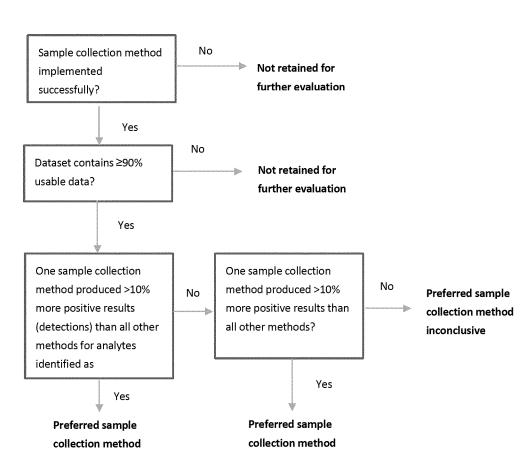


Figure 3-1: Phase I Evaluation Process Flow Chart

Notes:

Section 4 describes the results of the evaluation process with respect to implementability (Step 1). The results of the evaluation process with respect to analytical data evaluation (Steps 2 to 4) are described in Section 5. Results are documented on the comparison charts outlined in Worksheet #11 of the QAPP (Tierra 2013) (included as Appendices A to J) and referenced in the applicable sections(s) of this Phase I Report.

^{1.} Steps 1 and 2 were carried out individually for each analytical group, for each sampling method, and for each sampling even t and attempt.

^{2.} If for a given analytical group, no sample collection method produced greater than 10% more positive results (detections) than all other methods, then the preferred sample collection method for that analytical group was identified as inconclusive.

4. Implementation Evaluation

As discussed in Section 3, the first step in the evaluation process is an assessment of implementability. Implementability is defined as the degree to which each sample collection method was successful in collecting the required samples for laboratory analysis and meeting the minimum analytical SOP requirements as defined in the QAPP (Worksheets #19-1 through 19-4; Tierra 2013). For any given sampling attempt, if a sample collection method was not successful in collecting samples for laboratory analyses, it would not be considered for further evaluation and was not included in the comparison of sample collection methods for that analytical group(s).

The following sections discuss implementation challenges common to all sample collection methods for consideration during the ultimate selection of sample collection method(s). A comparison of the sampling approaches with respect to implementation challenges encountered and ability to successfully generate target mass/volume for laboratory analysis is presented below.

4.1 Implementation Requirements and Challenges

Mobilization requirements were common for all sample types. Specific mobilization requirements and challenges addressed during the sample collection activities included the following:

Ш	Site access and sidewalk closure and occupancy permit
	Coordination with Newark Police
	Weather monitoring
Ш	Coordination with PVSC
	Storm duration

A sidewalk closure and occupancy permit was obtained from the City of Newark to access and stage the sample collection system at the Clay Street CSO. Such permit would be required for any sampling approach utilized in Phase II. The permit application was initially prepared and approved prior to the first sample collection event and renewed every 30 days during the Phase I sampling program. Therefore, the permit was in place at all times during the potential sample collection period. Typically, the City of Newark does not issue permit renewals and requires submitting a new permit application. However, because the sample collection task is rainfall dependent, the City of Newark agreed to issue permit renewals every 30 days. Sampling location within differenttownships may be subject to different requirements.

Tierra coordinated with the City of Newark police during sample collection to provide traffic/site safety control in accordance with New Jersey Department of Transportation regulations. The Clay Street CSO sampling location is located at the intersection of Clay Street and McCarter Highway in Newark, New Jersey. Due to

Title: Phase I Evaluation/Recommendation Report

Revision Number: 2

Revision Data: June 2016

heavy traffic and the need to occupy the sidewalk, police support was required to provide traffic control. Additionally, site safety was needed to facilitate collection of bulk samples during nights and weekends.

Weather monitoring was performed during Phase I sample collection to determine an appropriate time to initiate mobilization for sample collection. The QAPP (Tierra 2013) states the following criterion for mobilization: "For a precipitation event to trigger mobilization for sample collection, the event must be anticipated to produce at least 0.2 inch of rain with an average intensity of at least 0.05 inch per hour with no more than 4 consecutive dry hours during the event." Based on the target storm duration of four to six hours for sample collection, the length of the rainfall period expected to meet the mobilization criteria was also considered. A four to six hour sample collection period was targeted as this was the length of time anticipated to be needed to collect enough solids within the CFC to obtain all samples based on the limited existing TSS data for CSO effluent. Tierra screened various weather forecast providers to select a precipitation forecast provider to predict storm events to prepare and quickly respond to potential storm events for sample collection. Given the capabilities of the weather services evaluated, The Weather Channel and Weather Underground were used for general, long-term (7- to 10-day) weather monitoring, while the National Oceanic and Atmospheric Administration's National Weather Service (NOAA's NWS) was used for more precise monitoring (6- and 3-day forecasts) to evaluate the potential precipitation on an hourly basis. The NOAA's NWS station located at the Newark Liberty International Airport, New Jersey was identified as the location closest to the CSO location for the Phase I CSO/SWO sampling program. During periods of anticipated sample collection, monitoring of the forecast weather from the three providers was reviewed on a daily basis. Tierra monitored the forecast daily and whether there were events within 10, 7, 6, or 3 days with the potential to trigger mobilization for sample collection. Tierra then notified other members of the project team if an event was identified to trigger mobilization.

Following the initiation of Phase I sample collection, based on a comparison of actual (hourly precipitation data in inches available through NOAA's NWS) and predicted precipitation data and overflows recorded at the Clay Street CSO for various storm events, the mobilization criterion was modified from an average rainfall intensity of at least 0.05 inch per hour to an average intensity of at least 0.03 inch per hour. It was identified that several overflow events were missed due to the 0.05 inch per hour average rainfall intensity mobilization criterion, and that an average intensity of 0.03 inch per hour resulted in sufficient overflow conditions at the Clay Street CSO. Therefore, the mobilization criterion was changed to 0.03 inch per hour for rainfall intensity. The mobilization criterion for total rainfall remained the same (0.2 inch of rain).

Although the modification to the mobilization criteria resulted in mitigating missed overflows, sample collection could not be completed during six mobilization events due to other factors, including the following:

No raintall or less than anticipated raintall, contrary to forecasted conditions
No overflow occurrence during rain events that met the mobilization criteria

Title: Phase I Evaluation/Recommendation Report

Revision Number: 2

Revision Data: June 2016

Overflow lasted for less than the target duration of 4 to 6 hours, resulting in no sample collection
Water level in the diversion chamber manhole was low (approximately1 feet from the bottom), limiting the ability of the intake tubing to pump effluent and remain 1 foot off the bottom as required by the QAPP (Tierra 2013)
An operational issue with the CFC.

During anticipated storm events, Tierra coordinated with PVSC regarding the timing of regulator gate valve openings at the sampling location. During a storm event, as soon as the regulator gate valve was opened at the Clay Street CSO, PVSC contacted Tierra to notify them of the gate opening and overflow conditions at the Clay Street CSO. Sample collection was initiated following PVSC confirmation regarding gate opening. Following the storm event, PVSC contacted Tierra with notification that the regulator gate valve was closed at the Clay Street CSO, indicating the end of overflow conditions. PVSC had informed Tierra that overflows can occur without the regulator gate being opened. During one mobilization event on October 7, 2013, the sampling crew observed overflow at the Clay Street CSO location and bulk sample collection was initiated, although Tierra did not receive notification that the regulator gate valve had been opened (and, therefore, presumably was not).

4.2 Evaluation of Sampling Methods

The following subsections discuss the challenges associated with each of the sampling methods (HSM, LSM, whole water, and grab metals) and the measures taken to address such challenges. The systematic evaluation of these methods is governed by the implementability of the sampling methods and the ability to generate target sample mass/volume to accommodate the full suite of target analytes.

4.2.1 High-Solids Mass

4.2.1.1 High-Solids Mass Particulate

As described in Section 2, HSM particulate samples were generated from the solids retained in the CFC bowl, and the samples were processed and shipped to analytical laboratories the day after bulk sample collection.

Implementation Challenges and Logistics

Minor challenges were encountered during sample collection, and modifications were implemented to address these challenges.

The CFC setup is more labor intensive as compared to the other sample collection methods (whole water and LSM). The CFC sampling equipment has moving parts and thus the potential for breakdown. To address the labor requirements and the complexity of operating the system, prior to the start of Phase I sample collection, an adequate number of personnel were trained to setup and operate the centrifuge and were required to be familiar with the SOPs and manufacturers' specifications of the multiple systems in the sample collection trailer. As part of the CSO/SWO investigation, a field demonstration and testing of the sample collection system was conducted on August 24, 2012 at the Ivy Street CSO outfall located in Kearny, New Jersey.

During all sampling attempts at the Clay Street CSO, two material types ("fines" and "non-fine paper-like material") were encountered in the CFC bowl during HSM particulate sample collection. The challenge was to create a homogeneous particulate sample for laboratory analyses. A modification to the SOP was implemented and a stainless steel blender was used to process and blend the fines and non-fines material to create a homogeneous particulate sample for laboratory analysis. SOP No. 4 – Sample Processing and Collection (Tierra 2013) provides additional details on the blending process. The HSM particulate placed into sample containers by the field team during the first attempt of the first event consisted of only the fines portion of the HSM particulate material. Because this sample was not homogenized with the non-fines portion of the particulate, as was the case during all subsequent sampling attempts and events, data from this first sampling attempt was not considered useable for purposes of the Phase I evaluation and were not considered further and are not included in this Phase I Report. PCDDs/PCDFs and PCB congener sample results for Event #1, Attempt #1 are included in Appendix A and B, respectively.

During pre-Phase I blank collection and decontamination activities, it was observed that small particulates remained in the CFC following prescribed decontamination procedures and caused potential issues with CFC operation. It was decided to add a decontamination step to power wash the CFC bowl to remove the residual particulates. The power-washing step adds more time to the decontamination process, but avoids potential operational issues with the CFC.

A significantly fewer number of sample containers were required to ship the HSM particulate samples (primary and contingency) compared to the LSM and whole water sample collection methods and, therefore, resulted in lower actual bottle breakage during shipping and required less time for sample packaging and shipment.

Ability to Generate Target Sample Mass/Volume

The HSM sample collection method generated sufficient solids mass required for the targeted sample analyses. A minimum of two sampling attempts was needed to generate the targeted solids mass (2,400 grams; including QA/QC samples and primary and contingency samples) during each sampling event. During a single sampling attempt (6-hour sample collection), sufficient solids mass (approximately 1,550

grams) was generated to collect primary samples (including QA/QC) to accommodate the full targeted analytical groups (1,130 grams). An additional sampling attempt was needed to accommodate contingency sample mass for laboratory analysis. Note that this observation is based on one sampling location (Clay Street CSO) and solids mass retained in the CFC will vary at different CSO locations as it is dependent on the influent TSS.

Contingency Mass/Volume

No contingency samples were used in the HSM particulate sample collection method (see Appendix C).

4.2.1.2 High-Solids Mass Dissolved

As described in Section 2, the HSM dissolved samples were generated by subsampling from the HSM dissolved bulk sample collection tank using a small-diameter peristaltic pump and dedicated Teflon® tubing, and the samples were processed and shipped to analytical laboratories the day after bulk sample collection

Implementation Challenges and Logistics

The challenges identified above for HSM particulate sampling with regards to operation and decontamination of the CFC apply to the HSM dissolved sampling.

A secondary tank was needed around the HSM bulk sample collection tank to facilitate the placement of ice which was used to immediately begin to chill, and to then maintain, the cool temperature of the HSM dissolved bulk sample.

Due to the high sample volume required for each analytical group, larger (than typically used for standard aqueous analytical methods) sample containers were required to ship HSM dissolved samples compared to the HSM particulate sampling method and, therefore, resulted in bottle breakage during shipping and required more time for sample processing and shipment. However, approximately the same number of sample containers were needed to collect the HSM dissolved samples as the LSM bulk and whole water samples. Additional sample packaging steps (e.g., bubble wrap, pre-cut foam inserts) were undertaken to mitigate bottle breakage during sample shipment.

Ability to Generate Target Sample Mass/Volume

One successful six-hour sampling attempt/event was needed to generate the target sample volume (approximately 230 liters; including QA/QC samples and primary and contingency samples) to accommodate the full target analytical groups. However, as noted in Section 2, only a portion of the effluent stream from the CFC was diverted to the HSM bulk sample collection tank. The rate at which the effluent

Title: Phase I Evaluation/Recommendation Report

Revision Number: 2

Revision Data: June 2016

was pumped from the CFC effluent stream into the HSM bulk sample collection tank could potentially be modified to collect the required volume for HSM dissolved samples within a shorter time period.

Contingency Mass/Volume

HSM dissolved contingency volumes utilized are described below and are outlined in Appendix C.

□ Event #1, Attempt #1 HSM dissolved: Two contingency bottles were utilized for PCB congener analyses due to breakage of primary sample containers observed upon laboratory receipt.

4.2.2 Low-Solids Mass

4.2.2.1 Low-Solids Mass Bulk Sample Collection

Similar to HSM dissolved samples, LSM bulk samples were generated for laboratory analyses by subsampling from the whole water/LSM bulk sample collection tank using a small-diameter peristaltic pump and dedicated Teflon® tubing, and the samples were processed and shipped to analytical laboratories the day after bulk sample collection. The laboratory completed filtration of the LSM bulk sample to generate LSM particulate and LSM dissolved samples.

Implementation Challenges and Logistics

The challenges identified above for HSM dissolved sampling (i.e., need for a secondary tank and large sample volumes/containers) apply to the LSM bulk sampling.

LSM bulk sample collection is similar to HSM dissolved sample collection, except the LSM bulk sample is collected prior to the CFC. As such, LSM bulk sample collection setup is generally less labor intensive compared to the HSM sample collection method.

As discussed in Section 2, the LSM/whole water bulk sample collection tank was double-lined with a Teflon® liner. During sample processing activities on December 9, 2013, a tear/rip was observed at the bottom of the inside Teflon® liner of the double-lined LSM bulk/whole water bulk sample collection tank after mixing and subsampling activities began. Water was collected from within the inner liner of the double-lined tank, and excess water remained in the tank at the end of sampling. It was not necessary to collect water from between the two Teflon® liners. The potential for liner tear/rip was identified during design of the sample collection system, and the bulk sample collection tanks were double-lined with Teflon® liners to avoid potential for bulk effluent to leak from the Teflon® liner and contact the tank. As such, no negative impacts to the sample were identified due to the identified tear/rip.

Revision Data: June 2016

Ability to Generate Target Sample Mass/Volume

One successful 6-hour sampling attempt/event was needed to generate the target sample volume (approximately 450 liters, including QA/QC samples and primary and contingency samples) to accommodate the full target analytical groups. However, as noted in Section 2, only a portion of the effluent stream from the manhole was diverted to the LSM bulk sample collection tank. The rate at which the effluent was pumped from the effluent stream into the LSM bulk sample collection tank could potentially be modified to collect the required volume for LSM bulk samples within a shorter time period.

4.2.2.2 Low-Solids Mass Bulk Laboratory Filtration

As described in Section 2, LSM bulk samples were generated by filtration at the laboratory.

Implementation Challenges and Logistics

The laboratory successfully filtered all of the LSM bulk samples using the primary approach. Although filtration of LSM bulk samples was relatively time consuming (as described below), the use of the secondary approach was not necessary.

The LSM bulk sample separation procedure is labor intensive due to the preparatory decontamination and setup requirements of the multi-component equipment. The LSM bulk sample separation equipment (for both the primary and secondary approach), comprise multiple components, including various tubing and filter media housing. These component parts require rigorous decontamination, and associated blank collection, between uses in separating LSM bulk material obtained from different sampling events.

Additionally, the filter media used to separate the LSM bulk samples is pre-cleaned in lots prior to use to verify that filters are not contributing any contamination to the LSM samples during bulk sample filtration. A representative filter from the lot is selected and submitted for laboratory analysis. Results of the analyses are used to certify that the filter media are contaminant-free or to establish background contaminant concentrations in the filter media as applicable. Pre-cleaned filter media must be re-certified to re-establish contaminant background concentration if not used to separate samples over a period greater than 6 months from the initial evaluation.

The LSM bulk sample separation procedure is time consuming as it requires the filtration of large volumes of LSM bulk sample to meet the analytical sensitivity requirements established in the QAPP (Tierra 2013). Table 4-1 below identifies the volume requirements for each analytical group.

Revision Data: June 2016

Table 4-1
LSM Bulk Liquid Volume Requirements by Analytical Group

Analytical Group	Minimum Sample Volume Required (liters)	Actual Sample Volume Collected per Event (liters)
PCDD/PCDFs	40	40
PCB Congeners	20	20
Organochlorine Pesticides	10	10
SVOCs	10	10
SVOC SIM	10	10
Aroclor PCBs	4	4
Chlorinated Herbicides	4	4
POC/DOC	16	16
TSS	3	3
TDS	1.5	1.5

Minimum sample volume requirements listed above are per event and include the primary sample, field duplicate, and associated QA/QC samples. During Phase I, approximately 120 liters of LSM bulk sample were collected and processed during each event requiring approximately 48 labor hours. This volume/time does not take into consideration contingency volume that might be needed.

Ability to Generate Target Sample Mass/Volume

The LSM bulk sample filtration process did generate acceptable target sample volume for LSM dissolved samples. However, the LSM bulk sample filtration process was insufficient in generating the target sample mass for LSM particulate samples. Table 4-2 provides the targeted and corresponding actual LSM bulk sample volume filtered to produce the LSM dissolved samples. Table 4-3 provides the targeted sample mass for LSM particulate samples for each analytical group per event, as well as the corresponding actual mass of LSM particulate samples collected and analyzed by the laboratory during Phase I.

Revision Data: June 2016

Table 4-2
Targeted LSM Dissolved Volume and Corresponding Actual LSM Bulk Volume Filtered by Analytical Group

Analytical Group	Targeted LSM Dissolved Sample Volume (liters) ^a	Event #1, Attempt #1 LSM Bulk Volume Filtered (liters) ^{b,c}	Event #1, Attempt #2 LSM Bulk Volume Filtered (liters) ^b	Event #1, Attempt #3 LSM Bulk Volume Filtered (liters) ^b	Event #2, Attempt #2 LSM Bulk Volume Filtered (liters) ^{b,d}
PCDD/PCDFs	10	10.035	-	9.663	9.476
PCB Congeners	5	4.957	-	5.009	4.819
Organochlorine Pesticides	2.5	_	2.558	-	2.430
SVOCs	2.5	_	2.363	-	2.418
SVOC SIM	2.5	_	2.530	-	2.400
Aroclor PCBs	1	-	0.979	-	1.013
Chlorinated Herbicides	1	_	0.984	1.053	1.042
POC/DOC	4	-	4.057	-	4.147

Notes:

- a. Target volume is for sample only and does not include QC volume requirements.
- b. LSM bulk filtered volume presented are that of the original field sample only (without additional QC volume requirements) allowing direct comparison with the target volume value provided for each analytical.
- c. As a result of only the "fine" material being analyzed for E vent #1, Attempt #1, PCDDs/PCDFs and PCB congener samples from Event #1, Attempt #1 were "replaced" by Event #1, Attempt #3. T herefore, Event #1, Attempt #1 results were not included as part of the data evaluation process.
- d. No LSM samples were collected during Event #2, Attempt #1.
- = analytical group was not analyzed

Table 4-3
Targeted LSM Particulate Mass and Corresponding Actual LSM Particulate Mass by Analytical Group

Analytical Group	Targeted LSM Particulate Mass (grams) ^a	Event #1, Attempt 1 LSM Particulate Mass (grams) ^b	Event #1, Attempt #2 LSM Particulate Mass (grams)b	Event #2, Attempt #2 LSM Particulate Mass (grams) ^b	Event #1, Attempt #3 LSM Particulate Mass (grams) ^b
PCDD/PCDFs	1.5	0.370°	-	0.079	0.077
PCB Congeners	0.75	0.183°	_	0.040	0.040
Organochlorine Pesticides	0.375	-	0.166	0.020	-
SVOCs	0.375	-	0.163	0.020	-
SVOC SIM	0.375	-	0.160	0.020	-
Aroclor PCBs	0.15	-	0.068	0.008	-
Chlorinated Herbicides	0.15	-	0.064	0.009	0.008
POC	0.60	-	0.263	0.010	-

Notes

- a. Target sample mass was based on a historical TSS average of 150 milligrams per liter (mg/L). These values reflect the minimum sample mass set as a requirement for a single sample analysis and do not include additional QC mass requirements.
- b. LSM particulate mass values observed during the field investigation are that of the original field sample only (without additional QC mass requirements) allowing direct comparison with the target mass value provided. LSM particulate samples were not collected during Event # 2, Attempt # 1.

Title: Phase I Evaluation/Recommendation Report

Revision Number: 2

Revision Data: June 2016

c. As a result of only the "fine" material being analyzed for E vent #1, Attempt #1, PCDDs/PCDFs and PCB congener samples from Event #1, Attempt #1 were "replaced" by Event #1, Attempt #3. Therefore, Event #1, Attempt #1 results were not included as part of the data evaluation process.

- = analytical group was not analyzed

The low mass obtained for the LSM particulate samples is related to significantly lower (as low as 8 mg/L) than anticipated (150 mg/L) TSS concentrations observed during the sampling events/attempts at the Clay Street CSO. Reduced sample mass has a direct relationship with reduced analytical sensitivity; however, the LSM sample results were retained for further evaluation as part of the Phase I evaluation process. The smaller than anticipated sample size obtained for LSM particulates may be linked to the larger number of non-detected results observed for many of the constituents of concern (COCs) as a direct cause and effect. This is especially true for the hydrophobic constituents, which are associated in large part with the particulate, rather than the dissolved-phase of the CSO overflow. This is a limitation of the LSM sample collection method. Even if the anticipated LSM particulate sample size had been collected, the mass of particulates obtained would have been approximately 10 to 100 times less than the HSM particulate sample mass. Therefore, it is unclear if the targeted LSM particulate sample size would have produced a greater number of positive results for COCs when compared to the HSM particulate samples.

To account for potential low TSS and corresponding low LSM particulate sample mass during future sampling events, the possible addition of real-time TSS monitoring using a turbidimeter or similar equipment will be evaluated to make field adjustments for the volume of water that needs to be collected for LSM bulk samples.

Contingency Mass/Volume

No contingency sample masses or volumes were used in the LSM sample collection method (see Appendix C).

4.2.3 Whole Water

As described in Section 2, whole water samples were generated for laboratory analyses by subsampling from the LSM/whole water bulk sample collection tank using a small-diameter peristaltic pump and dedicated Teflon® tubing, and the samples were processed and shipped to analytic all aboratories the day after bulk sample collection.

The whole water sampling method is identical to the LSM bulk sampling method, with the only difference being there is no laboratory filtration to generate particulate and dissolved samples.

Contingency Mass/Volume

Whole water contingency volumes utilized are described below and are outlined in Appendix C to this Phase I Report.

- Event #1, Attempt #1 Whole Water: Thirty-three contingency bottles were utilized for PCDD/PCDFs and PCB congener analyses due to breakage in the primary sample upon laboratory receipt and several coolers being out of temperature range. Further, in the case of PCDD/PCDFs analysis, the sample, matrix spike, and matrix spike duplicate were re-extracted using contingency volume after solid-phase extraction disc clogging problems occurred during the original extraction.
- □ Event #2, Attempt #2 Whole Water. Four contingency bottles were utilized for organochlorine pesticide analysis of the primary sample and duplicate sample due to the delayed sample arrival of the primary samples to the laboratory. The laboratory was instructed to only use contingency volumes for the entire analysis (i.e., primary sample, duplicate, matrix spike, and matrix spike duplicate).

Sixteen contingency bottles were utilized for PCDD/PCDFs analysis due to the delayed sample arrival of the primary and duplicate samples to the laboratory. The laboratory was instructed to only use contingency volumes for all analyses (i.e., primary sample, duplicate, matrix spike, and matrix spike duplicate).

Eight contingency bottles were utilized for PCB congener analysis due to the delayed sample arrival of the primary and duplicate samples to the laboratory. The laboratory was instructed to only use contingency volumes for all analyses (i.e., primary sample, duplicate, matrix spike, and matrix spike duplicate).

□ Event #1, Attempt #3 Whole Water: Four contingency bottles were utilized for PCDD/PCDFs analysis due to breakage of one of the four primary bottles for the primary sample. The laboratory was instructed to only use the contingency volumes for the sample analysis.

4.2.4 Grab Metals

As described in Section 2, samples for grab metals, including mercury and methyl mercury analyses, were collected directly from the effluent stream into sample containers and shipped within 24 hours (to meet holding time requirements) to the analytical laboratory for analysis.

Implementation Challenges and Logistics

No significant challenges were encountered during implementation of grab metals sampling. However, with regards to ease of implementation, adequate lead time (approximately 2 to 3 weeks) is required for the

laboratory to decontaminate tubing and sample containers in accordance with the trace metals sampling protocol (USEPA 1996). Additionally, CH and DH sampling procedures needed to be implemented in accordance with SOP No. 5 – Metals Sampling via Method 1669 Sampling Ambient Water for Trace Metals at EPA Water Quality Criteria Levels (USEPA 1996) (Tierra 2013). The CH and DH procedures require additional preparation and implementation time in the field. The samples for metals (total and dissolved) were not preserved in the field. To meet the analytical method holding time requirements, metals samples were processed and shipped via overnight carrier within 24 hous of sample collection.

Ability to Generate Target Sample Mass/Volume

The sampling method was able to generate the target sample volume during each sampling event for the full target analytical groups.

Contingency Mass/Volume

No contingency volumes were used in the grab metals collection (see Appendix C).

4.3 Summary of Implementability Evaluation

In summary, with the exception of the samples collected during Event #1, Attempt #1 (see Section 4.2.1.1), all three sampling approaches (HSM, LSM, and whole water) were successful in collecting the required field samples for laboratory analyses for all analytical groups during the sampling events/attempts at the Clay Street CSO. Therefore, all samples collected met the evaluation criteria based on implementability and were retained for further evaluation. However, as noted in Section 2, multiple attempts were needed to incrementally (following the analytical hierarchy established in the QAPP) complete the overall sample volume requirements and the LSM particulate samples did not meet the targeted sample mass.

5. Analytical Data Evaluation

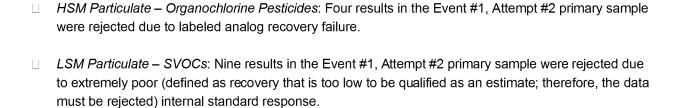
This section presents the results of Steps 2, 3, and 4 of the Phase I data evaluation process.

5.1 Data Usability

The second step of the evaluation process is an evaluation of the quality of the data generated. As stated above, validated data must contain a minimum of 90% usable data to be further assessed in the evaluation process. Table 5-1 below contains a summary of data that did not meet this criterion and, therefore, was not considered further in the evaluation process. Each is discussed in further detail below.

Table 5-1
Summary of Data Quality Failures

Sample Collection Method and Analytical Group	Event/ Attempt	Primary/ Duplicate Sample	Total Number of Results Reported	Number of Results Affected	% of Results Affected
HSM Particulate – Organochlorine Pesticides	Event #1, Attempt #2	primary	28	4	14
LSM Particulate – SVOCs	Event #1, Attempt #2	primary	50	9	18
HSM Dissolved – SVOCs	Event #1, Attempt #2	primary	50	8	16
HSM Dissolved – SVOCs	Event #1, Attempt #2	duplicate	50	8	16
HSM Particulate – VOCs	Event #1, Attempt #2	primary (fines)	6	4	67
HSM Particulate – VOCs	Event #1, Attempt #2	primary (non-fines)	6	4	67
HSM Particulate – VOCs	Event #1, Attempt #2	duplicate (fines)	6	4	67
HSM Particulate – VOCs	Event #2, Attempt #1	primary (fines)	6	4	67
HSM Particulate – VOCs	Event #2, Attempt #1	primary (non-fines)	6	5	83
HSM Particulate – VOCs	Event #2, Attempt #1	duplicate (fines)	6	4	67



Revision Data: June 2016

Ш	HSM Dissolved – SVOCs: Sixteen results in the Event #1, Attempt #2 primary and duplicate samples
	were rejected due to extremely poor (defined as recovery that is too low to be qualified as an estimate;
	therefore, the data must be rejected) internal standard response.

HSM Particulate – VOCs: Twenty-five results in the Event #1, Attempt #2 and Event #2, Attempt #1 primary (fines), primary (non-fines), and duplicate (fines) samples were rejected due to low internal standard responses.

Note that these data quality issues were related to laboratory performance and are not likely sample collection technique dependent.

All other data for each sampling method and analytical group met the usability requirements set out in the QAPP (Tierra 2013) and were considered further in the evaluation process.

5.2 Decontamination

As discussed in Section 2.4, applicable decontamination procedures were applied throughout the Phase I sample collection program in accordance with SOP No. 6 – Decontamination (Tierra 2013). Between sampling events, a full decontamination of the sample collection system was performed in accordance with Section 2.2.2 of SOP No. 6: Decontamination, included in the QAPP (Tierra 2013). Field, rinsate and equipment blanks were collected in accordance with Section 2.4 of SOP No. 6: Decontamination. Positive results identified in the field, rinsate, and equipment blanks collected during Phase I, and associated field blank implications on the data evaluation process are described in Section 5.3.

5.3 Field Blank Results and Affected Sample Results

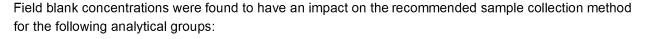
During the data validation process, positive sample results associated with analytes identified in a field blank were assessed per USEPA Region 2 and other data validation guidance provided in the approved QAPP (Tierra 2013). Positive sample results that fell within the affected concentration range as defined in the validation guidance, were qualified "U", not detected. The number of positive sample results qualified as "U" based on field blank contamination overall are included in Appendix D.

Tierra assessed the potential impact of field blank concentrations on the conclusions of the recommended sample collection method. The details of this assessment are included in Appendix E. The following assumption was made in order to assess the potential impact of field blank concentrations. For the purpose of this evaluation, all detected results as reported by the laboratory prior to validation, are assumed to be those of compounds present in the field sample collected, and not artifacts of background concentrations.

Title: Phase I Evaluation/Recommendation Report

Revision Number: 2

Revision Data: June 2016



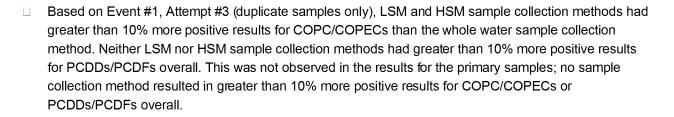
	PCB Congeners - Event #2, Attempt #2 (primary sample)
***************************************	Organochlorine Pesticides - Event #1, Attempt #2 (duplicate sample) and Event #2, Attempt #2 (primary sample)
	SVOCs SIM - Event #1, Attempt #2 (primary sample) and Event #1, Attempt #2 (duplicate sample)
and the de	Chlorinated Herbicides - Event #2, Attempt #2 (primary sample), Event #2, Attempt #2 (duplicate sample), Event #1, Attempt #3 (primary sample), and Event #1, Attempt #3 (duplicate sample).

5.4 Steps 3 and 4: Frequency of Detections

Data for a given analytical group and sampling method that were not eliminated from the evaluation process during Steps 1 or 2 were assessed in Steps 3 and 4 based on frequency of detections as defined above. A summary of the Steps 3 and 4 evaluations per analytical group are summarized below. In addition, a summary of the overall result of the evaluation process is also provided. As discussed in Section 4, the HSM particulate placed into sample containers by the field team during the first attempt of the first event consisted of only the fines portion of the HSM particulate material. Because this sample was not homogenized with the non-fines portion of the particulate, as was the case during all subsequent sampling attempts and events, data from this first sampling attempt was not considered useable for purposes of the Phase I data evaluation.

5.4.1 Polychlorinated Dibenzo-p-dioxins/Polychlorinated Dibenzofurans

All three sample collection and processing methods (LSM, HSM, and whole water) were evaluated for the PCDD/PCDFs analytical group. Samples (primary sample and field duplicate) were collected for PCDD/PCDF analysis during Event #1, Attempt #3 and Event #2, Attempt #2. A summary of the findings of the evaluation Steps 3 and 4 for PCDD/PCDF data are provided below. Detailed evaluation sheets (Worksheet #11) can be found in Appendix F.



□ Based on Event #2, Attempt #2 (primary and duplicate samples), the HSM sample collection method had greater than 10% more positive results for COPC/COPECs than the LSM and whole water sample collection methods.

Overall, the recommended sample collection method(s), if any, based on the results of the Phase I evaluation criteria (Steps 1 to 4) for PCDDs/PCDFs is summarized in Table 5-2 below.

Table 5-2
Recommended Sample Collection Method – PCDDs/PCDFs

	Event #1, Attempt #3	Event #2, Attempt #2
Primary Sample	Inconclusive	HSM
Duplicate Sample	LSM/HSM	HSM

5.4.2 Polychlorinated Biphenyl Congeners

All three sample collection and processing methods (LSM, HSM, and whole water) were evaluated for the PCB congeners analytical group. Samples were collected for PCB congener analysis during Event #1, Attempt #3 and Event #2, Attempt #2. A summary of the findings of evaluation Steps 3 and 4 for PCB congener data are provided below. The detailed evaluation sheets (Worksheet #11) can be found in Appendix G.

- Based on Event #1, Attempt #3 (duplicate samples), the HSM sample collection method had greater than 10% more positive results for COPC/COPECs than the LSM and whole water sample collection methods. The results for the primary sample showed both HSM and LSM sample collection methods had greater than 10% more positive results for COPC/COPECs than the whole water sample collection method; however, the HSM sample collection method also had greater than 10% more positive results for PCB congeners overall.
- Based on Event #2, Attempt #2 (primary samples), the HSM sample collection method had greater than 10% more positive results for COPC/COPECs than the LSM and whole water sample collection methods. The results for the duplicate samples showed both HSM and LSM sample collection methods had greater than 10% more positive results for COPC/COPECs than the whole water sample collection method; however, the HSM sample collection method also had greater than 10% more positive results for PCB congeners overall.

Overall, the recommended sample collection method(s), if any, based on the results of the Phase I evaluation criteria (Steps 1 to 4) for PCB congeners is summarized in Table 5-3 below.

Revision Data: June 2016

Table 5-3
Recommended Sample Collection Method – PCB Congeners

	Event #1, Attempt #3	Event #2, Attempt #2
Primary Sample	HSM	HSM
Duplicate Sample	HSM	HSM

5.4.3 Aroclor Polychlorinated Biphenyls

All three sample collection and processing methods (LSM, HSM, and whole water) were evaluated for the Aroclor PCBs analytical group. Samples were collected for Aroclor PCB analysis during Event #1, Attempt #2 and Event #2, Attempt #2. A summary of the findings of evaluation Steps 3 and 4 for Aroclor PCB data are provided below. The detailed evaluation sheets (Worksheet #11) can be found in Appendix H.

- Based on Event #1, Attempt #2 (primary and duplicate samples), the HSM sample collection methods had greater than 10% more positive results for COPC/COPECs than the LSM and whole water sample collection methods.
- Based on Event #2, Attempt #2 (duplicate samples), the HSM sample collection method had greater than 10% more positive results for COPC/COPECs than the LSM and whole water sample collection methods. This was not observed in the results for the primary samples; no sample collection method resulted in greater than 10% more positive results for COPC/COPECs or Aroclor PCBs overall.

Overall, the recommended sample collection method(s), if any, based on the results of the Phase I evaluation criteria (Steps 1 to 4) for Aroclor PCBs is summarized in Table 5-4 below.

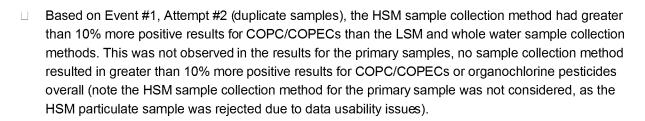
Table 5-4
Recommended Sample Collection Method – Aroclor PCBs

	Event #1, Attempt #2	Event #2, Attempt #2
Primary Sample	HSM	Inconclusive
Duplicate Sample	HSM	HSM

5.4.4 Organochlorine Pesticides

All three sample collection and processing methods (LSM, HSM, and whole water) were evaluated for the organochlorine pesticide analytical group. Samples were collected for organochlorine pesticides analysis during Event #1, Attempt #2 and Event #2, Attempt #2. A summary of the findings of evaluation Steps 3 and 4 for organochlorine pesticide data is provided below. The detailed evaluation sheets (Worksheet #11) can be found in Appendix I.

Revision Data: June 2016



Based on Event #2, Attempt #2 (primary samples), the HSM sample collection method had greater than 10% more positive results for COPC/COPECs than the LSM and whole water sample collection methods. This was not observed in the results for the duplicate samples; no sample collection method resulted in greater than 10% more positive results for COPC/COPECs or organochlorine pesticides overall.

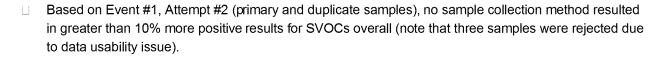
Overall, the recommended sample collection method(s), if any, based on the results of the Phase I evaluation criteria (Steps 1 to 4) for organochlorine pesticides is summarized in Table 5-5 below.

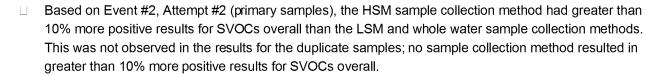
Table 5-5
Recommended Sample Collection Method – Organochlorine Pesticides

	Event #1, Attempt #2	Event #2, Attempt #2
Primary Sample	Inconclusive	HSM
Duplicate Sample	l HSM	Inconclusive

5.4.5 Semivolatile Organic Compounds

All three sample collection and processing methods (LSM, HSM, and whole water) were evaluated for the SVOC analytical group. Samples were collected for SVOC analysis during Event #1, Attempt #2 and Event #2, Attempt #2. A summary of the findings of evaluation Steps 3 and 4 for SVOC data are provided below. Note there are no COPECs that are SVOCs. The detailed evaluation sheets (Worksheet #11) can be found in Appendix J.





Overall, the recommended sample collection method(s), if any, based on the results of the Phase I evaluation criteria (Steps 1 to 4) for SVOCs is summarized in Table 5-6 below.

Table 5-6
Recommended Sample Collection Method – SVOCs

	Event #1, Attempt #2	Event #2, Attempt #2
Primary Sample	Inconclusive	HSM
Duplicate Sample	Inconclusive	Inconclusive

5.4.6 Semivolatile Organic Compounds Select Ion Monitoring

All three sample collection and processing methods (LSM, HSM, and whole water) were evaluated for the SVOC SIM analytical group. Samples were collected for SVOC SIM analysis during Event #1, Attempt #2 and Event #2, Attempt #2. A summary of the findings of evaluation Steps 3 and 4 for SVOC SIM data are provided below. The detailed evaluation sheets (Worksheet #11) can be found in Appendix K.

- Based on Event #1, Attempt #2 (primary and duplicate samples), the HSM sample collection method had greater than 10% more positive results for COPC/COPECs than the LSM and whole water sample collection methods. HSM sample collection method had greater than 10% more positive results for SVOC SIM overall.
- Based on Event #2, Attempt #2 (primary and duplicate samples), the HSM sample collection method had greater than 10% more positive results for COPC/COPECs than the whole water sample collection method but less than 10% more positive results for COPC/COPECs than the LSM sample collection method. Neither LSM nor HSM sample collection method had greater than 10% more positive results for SVOC SIM overall. These observations resulted in the LSM/HSM sample collection methods ranked as equivalent for the primary sample. This was not observed in the results for the duplicate sample. No sample collection method resulted in greater than 10% more positive results for COPC/COPECs or SVOCs SIM overall.

Overall, the recommended sample collection method(s), if any, based on the results of the Phase I evaluation criteria (Steps 1 to 4) for SVOCs SIM is summarized in Table 5-7 below.

Table 5-7
Recommended Sample Collection Method – SVOCs SIM

-	Event #1, Attempt #2	Event #2, Attempt #2
Primary Sample	HSM	LSM/HSM
Duplicate Sample	HSM	Inconclusive

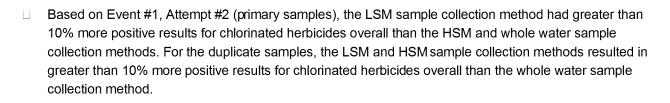
Revision Number: 2

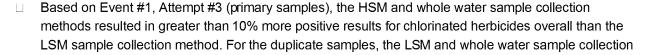
Revision Data: June 2016

5.4.7 Chlorinated Herbicides

All three sample collection and processing methods (LSM, HSM, and whole water) were evaluated for the chlorinated herbicides analytical group. Samples were collected for chlorinated herbicide analysis during Event #1, Attempt #2; Event #1, Attempt #3; and Event #2, Attempt #2. Three sets of samples were collected due to a laboratory error identified during the herbicide analysis of the HSM particulate sample from Event #2, Attempt #2. The HSM particulate herbicide results indicated that a laboratory control sample associated with the herbicide data had failed. In an attempt to produce results that would be free of qualification, the laboratory was asked to re-extract and re-analyze the sample. The laboratory reported that the remaining HSM particulate sample had developed a mold growth on the surface of the sample. It was decided that the presence of this mold could pose data quality issues; therefore, it was suggested to the USEPA that additional chlorinated herbicide samples be collected during the next sampling event (Event #1, Attempt #3). This was approved by the USEPA in an email correspondence on February 20, 2014 (USEPA 2014). Data from all three sampling events/attempts, including herbicide results from Event #2, Attempt #2 affected by the failed laboratory control sample, have been used in this evaluation. A summary of the findings of evaluation Steps 3 and 4 for chlorinated herbicides data are provided below. Note there are no COPECs that are chlorinated herbicides. The detailed evaluation sheets (Worksheet #11) can be found in Appendix L.

It should be noted that many of the positive chlorinated herbicide results were qualified as tentatively identified at an estimated concentration (NJ). This is a reflection of a larger than acceptable level of uncertainty as to both the qualitative identification of the analyte and the numerical value reported. Across all sample types collected during the three sampling events/attempts, 29 positive chlorinated herbicide results were reported. Of those 29 positive results, 16 were assigned an "NJ" flag during validation. A significant component of the data evaluation process is a comparison of the number of positive results reported between sample collection methods (Steps 3 and 4). Therefore, the conclusions of the data evaluation process, and thereby the selection of a recommended sample collection method, may have been impacted by the larger than acceptable uncertainty in qualitative analyte identification noted during herbicide data validation.





Revision Number: 2

Revision Data: June 2016

methods resulted in greater than 10% more positive results for chlorinated herbicides overall than the HSM sample collection method.

Based on Event #2, Attempt #2 (primary samples), the HSM sample collection method resulted in greater than 10% more positive results for chlorinated herbicides overall than the LSM and whole water sample collection methods. For the duplicate samples, the LSM sample collection method resulted in greater than 10% more positive results for chlorinated herbicides overall than the HSM and whole water sample collection methods.

Overall, the recommended sample collection method(s), if any, based on the results of the Phase I evaluation criteria (Steps 1 to 4) for chlorinated herbicides is summarized in Table 5-8 below.

Table 5-8
Recommended Sample Collection Method – Chlorinated Herbicides

	Event #1, Attempt #2	Event #1, Attempt #3	Event #2, Attempt #2
Primary Sample	LSM	HSM/whole water	HSM
Duplicate Sample	LSM/HSM	LSM/whole water	LSM

5.4.8 Cyanide

As per the QAPP (Tierra 2013), only HSM and whole water sample collection methods were evaluated for the cyanide analytical group since only whole water sample collection (and not LSM sample collection) were included in the CSO/SWO S&AP (USEPA 2008).

Samples were collected for cyanide analysis during Event #1, Attempt #2 and Event #2, Attempt #2. A summary of the findings of evaluation Steps 3 and 4 for cyanide data are provided below. Note cyanide is not a COPEC. The detailed evaluation sheets (Worksheet #11) can be found in Appendix M.

Based on Event #1, Attempt #2 and Event #2, Attempt #2 (primary and duplicate samples), cyanide data exhibited positive results for the analyte in the samples collected using HSM and whole water sample collection methods. Because cyanide is a single-component analytical group with 100% detections for both methods, one sample collection method did not produce greater than 10% more positive results (detections) than all other methods. Therefore, the recommended sample collection method(s) based on the Phase I evaluation criteria is inconclusive.

5.4.9 Volatile Organic Compounds

As per the QAPP (Tierra 2013), only whole water and HSM sample collection and processing methods were evaluated for the VOC analytical group since only whole water sample collection (and not LSM sample

Revision Number: 2

Revision Data: June 2016

collection) were included in the CSO/SWO S&AP (USEPA 2008). Samples were collected for VOC analysis during Event #1, Attempt #2 and Event #2, Attempt #1. However, samples collected using the HSM sample collection method were rejected due to data usability issues. Therefore, only data for samples collected via the whole water samples collection method were considered usable. The detailed evaluation sheets (Worksheet #11) can be found in Appendix N.

The whole water sample collection method was not selected as the recommended method for VOCs. A limited dataset was available to complete the data comparison between sampling approaches, and only data for samples collected via the whole water method were considered usable. Additional investigation is recommended during Phase II to evaluate sampling approaches for VOCs.

5.4.10 Total Extractable Petroleum Hydrocarbons

As per the QAPP (Tierra 2013), only whole water and HSM sample collection and processing methods were evaluated for the TEPH analytical group since only whole water sample collection (and not LSM sample collection) were included in the CSO/SWO S&AP (USEPA 2008). Samples were collected for TEPH analysis during Event #1, Attempt #2 and Event #2, Attempt #2. A summary of the findings of evaluation Steps 3 and 4 for TEPH data are provided below. Note TEPH is not a COPEC. The detailed evaluation sheets (Worksheet #11) can be found in Appendix O.

Based on Event #1, Attempt #2 and Event #2, Attempt #2 (primary and duplicate samples), TEPH data exhibited positive results for the analyte in the samples collected using both the HSM and whole water sample collection methods. Because TEPH is a single-component analytical group with 100% detections for both methods, one sample collection method did not produce greater than 10% more positive results (detections) than all other methods. Therefore, the recommended sample collection method(s) based on the Phase I evaluation criteria is inconclusive.

5.5 Impacts of Achieved Analytical Sensitivity

Sensitivity is related to the ability to compare analytical results with project quantitation limits (PQLs). Analytical detection limits should be at or below the PQLs to allow effective comparisons. All sample analytical results reported during Phase I of the CSO/SWO investigation were evaluated to determine if adequate sensitivity was achieved. The results for each analyte were cross-checked against the PQLs presented in Worksheet #15 of the QAPP (Tierra 2013). The results of this detailed evaluation are presented in the CSO/SWO Investigation Phase I Data Quality Usability Assessment Report (DQUAR; Tierra 2016). The DQUAR (Tierra 2016) is included as Appendix P.

The observation that data obtained for a particular sample type/collection method failed to meet established PQLs for specific analytical groups may have impacted the number of positive results

identified in those samples, thereby potentially impacting the data evaluation process. Tierra performed an evaluation of instances where PQL exceedances were identified to assess any potential impact on the data evaluation process and sample collection method selection. The results of this additional evaluation is also included in the DQUAR (Tierra 2016).

The following table summarizes the conclusions following assessment of the potential impact of PQL exceedances for each sample collection method during the data evaluation and selection process.

Table 5-9 Impact of PQL Exceedances

	PQL Exceedances May Have Impacted the Sample Collection Evaluation Process Yes/No													
Analytical Group	Whole Water	LSM Dissolved	LSM Particulate	HSM Dissolved	HSM Particulate									
PCDDs/PCDFs	No	NA	NA	NA	Yes									
PCB Congeners	Yes	Yes	Yes	Yes	No									
Organochlorine Pesticides	No	No	Yes	No	No									
SVOCs SIM	No	Yes	Yes	NA	Yes									
SVOCs	Yes	No	Yes	Yes	Yes									
Aroclor PCBs	NA	NA	Yes	NA	No									
Chlorinated NA Herbicides		NA	Yes	Yes	NA									
VOCs	NA	NA	NA	NA	No									

Notes:

NA= not applicable since non-detected results were not reported when or if PQL exceedances were noted for an analytical group.

5.6 Additional Data Evaluation

A side-by-side comparison of the HSM and LSM particulate and dissolved-phase concentrations and whole water was completed outside the scope of the data evaluation criteria as defined in the QAPP (Tierra 2013). Additionally based on comments received from the USEPA dated October 6, 2015 on this Phase I Report (Revision 0), and based on the results obtained for the Phase I sampling program, additional data evaluation was completed for select analytical groups to calculate summary statistics, compare results/concentrations, and evaluate trends to assist with development of the Phase II sampling program. Additional data evaluation was completed for the following analytical groups:

PCDD/PCDFS
PCB congeners

Title: Phase I Evaluation/Recommendation Report

Revision Number: 2

Revision Data: June 2016

,	Ossas a shi a sina a sa shi si da a
_	Organochlorine pesticides
	SVOCs
	SVOCs SIM
	Aroclor PCBs
	Chlorinated herbicides
	VOCs
	Cyanide
	TEPH

Findings and results of the additional data evaluation is included in Attachment 1 - Phase I Report Addendum – Additional Data Evaluation.

Revision Data: June 2016

6. Conclusion/Recommendation

Based on the Phase I evaluation process, the recommended sample collection methods per analytical group are identified below in Table 6-1. The HSM sample collection method is the preferred approach for certain hydrophobic contaminants, such as PCDDs/PCDFs, PCB congeners, Aroclor PCBs, and organochlorine pesticides. For PCB congeners, HSM was the recommended sample collection method for each sample collected (primary and duplicate) based on the Phase I evaluation process. For PCDDs/PCDFs, Aroclor PCBs, and organochlorine pesticides, HSM was the recommended sample collection method for half or more of the samples collected (primary and duplicate) based on the Phase I evaluation process. A preferred sample collection method for the remaining analytical groups was not definitive.

Table 6-1
Phase I Sample Collection Method Recommendations

Sample Collection Technique	PCDD/ PCDF	PCB Congeners	Aroclor PCBs	Organochlorine Pesticides	SVOCSSIM	SVOC	Chlorinated Herbicides	Cyanide	VOC	ТЕРН
LSM										
HSM						o	0	0	0	0
Whole Water										

Notes:

□ □ = selected sampling method

O = recommended sample collection method inconclusive

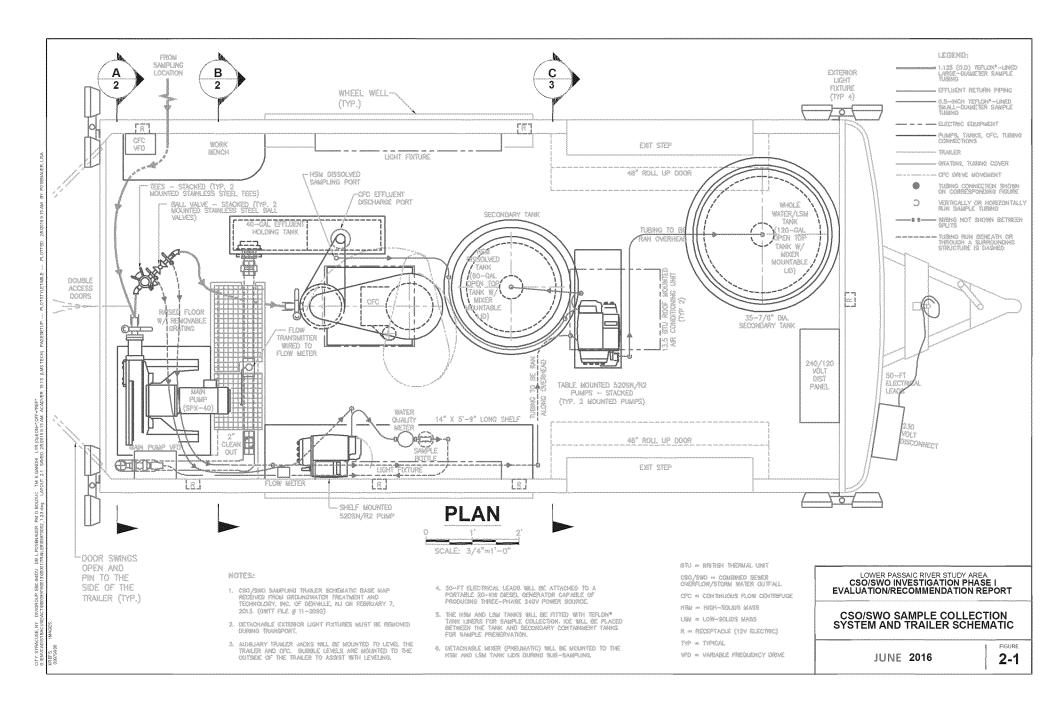
Based on the results of the Phase I evaluation discussed in this Phase I Report, it is recommended that a hybrid sample collection program be implemented for Phase II. Such a hybrid approach would focus on using the most appropriate sampling technique for each applicable parameter group. It is also recommended that Phase II be implemented in additional phases to continue to collect data and make adjustments (if needed) to meet program objectives. Given the number of additional sampling locations remaining to be sampled (eight CSOs, 10 SWOs, and one POTW sample [quarterly basis for 1 year]) during Phase II, an iterative evaluation of the Phase II data will allow flexibility in making adjustments to the program and help avoid collection of a large amount of data that do not meet program objectives.

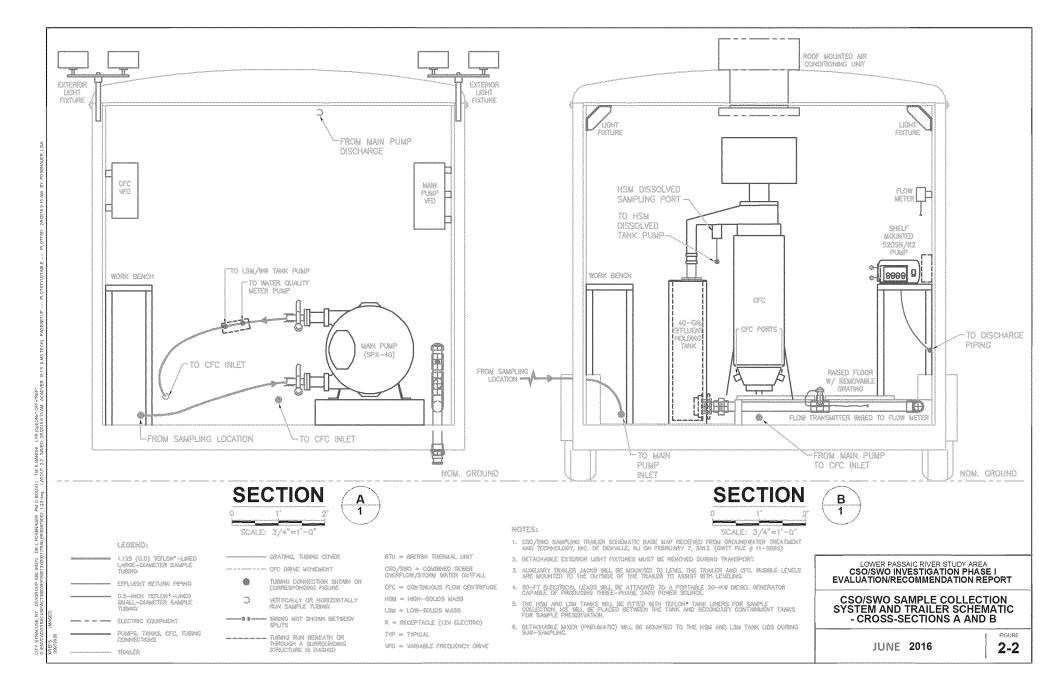
Tierra recommends a meeting with the USEPA to review the results of the Phase I evaluation and develop the approach and scope for the Phase II CSO/SWO investigation program that considers factors, including sampling technique, implementability, data needs, locations, and schedule.

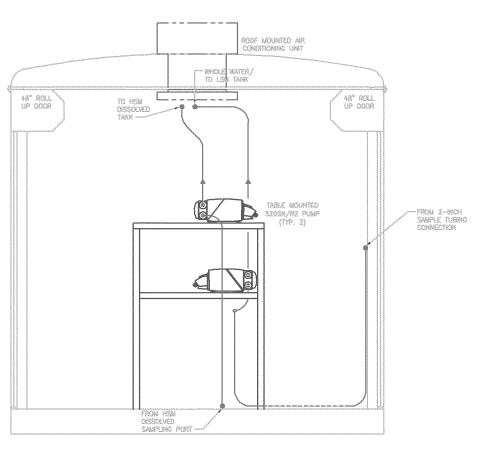
7. References

- Great Lakes Environmental Center. 2008. New York-New Jersey Harbor Estuary Program Contaminant Assessment and Reduction Program. New Jersey Toxics Reduction Work Plan Study I-G Project Report, February 2008.
- Malcolm Pirnie, Inc. 2008. Rain Event Program Narrative, Lower Passaic River Restoration Project (version 11/05/2008) Source: www.ourPassaic.org.
- The Louis Berger Group (in conjunction with Battelle HDR/HydroQual). 2014. Lower Eight Miles of the Lower Passaic River. Focused Feasibility Report. For U. S. Environmental Protection Agency, Region 2 and U.S. Army Corps of Engineers, Kansas City District.
- Tierra. 2002. Remedial Investigation Combined Sewer Overflow Investigation, Volume 1, Work Plan/Field Sampling Plan. May.
- Tierra. 2013. Combined Sewer Overflow/Stormwater Outfall Investigation Quality Assurance Project Plan. Lower Passaic River Study Area. Revision 3. September 2013.
- Tierra 2016. Combined Sewer Overflow/Stormwater Outfall Investigation Phase I Data Quality Usability Assessment Report, Revision 1. March 2016.
- USEPA. 1996. Method 1669, Sampling Ambient Water for Trace Metals at EPA Water Criterion Levels, U.S. Environmental Protection Agency, Office of Water Engineering and Analysis Division (4303), July 1996.
- USEPA. 2008. Combined Sewer Overflow/Stormwater Overflow Sampling and Analytical Plan, Revision No. 2.0. August.
- USEPA. 2014. Email Correspondence approving additional chlorinated herbicide samples. February 20.

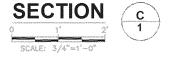
Figures







NOM. GROUND



DIVIGROUP EBC-IMDV DB L-POSBVAUER PM D.BOLDUC TM K.GANDH! LYR.(OptON="OFF="PEF" ACTB00099790013000011TRALER/09979002_1-23-54wg LAYOUT: 2.3 SAVED: 25.2016-9-15-AM ACADV

LEGENO:

*** 1-BOOK TEFLONF-LINED SAMPLE TURNING

EFFLUENT PICTURN FIFTIG

- 0.5-BYCH TEPLOHP-LINED SAMPLE TUBBRO

--- ELECTRIC EQUIPMENT

- PUMPS, TANKS, CFC, TUBBING CONNECTIONS

TRAILER

---- OFC DAVE MOVEMENT

TUBING CONNECTION SHOWN ON CORRESPONDING FIGURE

DHIBUT EFERS HUR YEARTHOEFECH NO YEEKSEW

--- WATERS NOT SHOWN BETWEEN SPLITS

THE THE TENDERS SENERTH OF THEOLOGY A SUPERCLUSIC STRUCTURE IS DARKED.

BITU = BRUTISH THERWAL LINET

CSO/SWO = COMMINED SEVER OVERFLOW/STORM WATER OUTFALL

OFC = CONTINUENTS FLOW CENTRIFUGE

HSM = HSGH-SOLIDS MASS

USH - LOW-SCLOS MASS

R - RECEPTACLE (12V ELECTRIC)

TWF = TYPHCAL

VFD = VARIABLE PREDIENCY DRIVE

NOTES:

- CSD/SMC BAMPLING TRAKER DEFENIATE BASE MAP RECEIVED FROM GROWNISK TREATMENT AND TED-MOLDST, NG. OF DERVALE, NJ ON FERSMARY 7, 2013. (ONTY FRE # 11-2082)
- 2. DETACHABLE EXTERIOR LIGHT FIXTURES MUST BE REMOVED DURING TRANSPORT.
- I AUGUSAY TRACER LACKS BUT SE MOUNTED TO LEVEL TWE TRACER AND CITC SUBSILE LEVELS AND MOUNTED TO THE OUTSIDE OF THE TRACER TO ASSIST WIN LEVELING.
- 4. SO-FT ELECTRICAL LEADS WELL BE ATTACHED TO A PERTABLE 20-RW DESEL ODERFATOR CAPAGE OF PROCEDURE THREE-PHASE 242V POWER SOURCE.
- 5. THE HEM AND LIM TARRE WILL BE PITTED WITH TEPLOR*
 TARK LIMERS FOR SAMPLE COLLECTION, ICE WILL BE PLACED
 BETTELD HE AND AND SECONDARY CONTAINERS FOR SAMPLE PRESENTATION.
- 4. DETACHABLE MINER (PHELMARC) WILL BE MICHITED TO THE HOM AND LISE TANK LOS DURING SUB-SAMPLING.

LOWER PASSAIC RIVER STUDY AREA CSO/SWO INVESTIGATION PHASE I EVALUATION/RECOMMENDATION REPORT

CSO/SWO SAMPLE COLLECTION SYSTEM AND TRAILER SCHEMATIC - CROSS-SECTION C

JUNE 2016

FIGURE **2-3**

STAINLESS FLOW ORIENTATION SEE NOT TO SCALE SHEP PLANE 8 (MINIMUM) X AHOM METALS COLLECTION TUBING SCHEMATIC OF WEIGHTED ROD/TUBING ASSEMBLY LOWER PASSAIC RIVER STUDY AREA CSO/SWO INVESTIGATION PHASE I EVALUATION / RECOMMENDATION REPORT SAMPLE NOTES: HIGH SOLIDS WASS ACRONYMS: FIGURE FOR VISUAL AID ONLY. EQUIPMENT SIZES ARE JUNE COLLECTION TUBING LOW SOLIDS MASS 2016 SAMPLE APPROXIMATE. N L

Appendix A

Event #1, Attempt #1 Results - PCDDs/PCDFs

EVENT 1 ORIGINAL SAMPLE - DIOXIN PR1CSOCLY**-01A

Positive Target Analyte Identification and Concentration Comparison^a

Analyte Identified	PR1CSOCLYWW-01A Whole Water ^b (pg/L)	LQʻ	VQ	PR1CSOCLYLD-01A LSM Dissolved ^f (pg/L)	LQ°	۷Q	PR1CSOCLYHD-01A HSM Dissolved ^b (pg/L)	ια°	VQ	% RPD	PR1CSOCLYLP-01A LSM Particulate ^b (pg/g)	LQ°	VQ	PR1CSOCLYHP-01A HSM Particulate ^b (pg/g)	LQ°	VQ	% RPD
2,3,7,8-TCDD				1995										2.36		j	
1,2,3,7,8-PeCDD							0.182	G			6.50	G		4.34	G	М	40
1,2,3,4,7,8-HxCDD	0.859	G					0.347	G			12.0	G		5.96		М	67
1,2,3,6,7,8-HxCDD	2.37	G		0.304	G	10	1.19	G		119	43.9	G		21.4		М	69
1,2,3,7,8,9-HxCDD	1.56	G		0.254	G	J	0.894	G		111	25.4	G		15.3		M	50
1,2,3,4,6,7,8-HpCDD	62.1		J	6.33		J	32.6		J	135	1940		J	672		М	97
OCDD	715		J	41.7		J	365		J	159	15700		J	9480	D	Μ	49
2,3,7,8-TCDF							0.148	G			5.49	G		4.76		М	14
1,2,3,7,8-PeCDF	0.228	G									4.09	G		3.76	G	N	8
2,3,4,7,8-PeCDF				0.0854	G		0.266	G		103	14.2	G		4.76	G	M	100
1,2,3,4,7,8-HxCDF	1.68	G		0.314	G	J	0.982	G		103	23.4	G		20.9		J	11
1,2,3,6,7,8-HxCDF				0.361	G	J	0.962	G		91	24.4	G		15.4		N	45
2,3,4,6,7,8-HxCDF	1.69	G		0.277	G	J	1.04	G		116	28.2	G		19.0		M	39
1,2,3,7,8,9-HxCDF											4.84	G		1.53	G	M	104
1,2,3,4,6,7,8-HpCDF	18.0			3.40		J	16.6			132	396		J	245		М	47
1,2,3,4,7,8,9-HpCDF	1.56	G									28.1	G		16.4		М	53
OCDF	36.6			6.05		J	37.0		J	144	790		J	486		М	48

COPCs/COPEcs listed in the FFS: 2,3,7,8-TCDD; 1,2,3,7,8-PeCDD; 1,2,3,6,7,8-HxCDD; 1,2,3,4,7,8-HxCDD; 1,2,3,7,8,9-HxCDD; 1,2,3,4,6,7,8-HpCDD; OCDD; 2,3,7,8-TCDF; 1,2,3,7,8-PeCDF; 2,3,4,7,8-PeCDF; 1,2,3,4,7,8-HxCDF; 1,2,3,4,

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

FFS = focused fesability study HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCDD/PCDF = polychlorinated dibenzo-p-dioxin/polychlorinated dibenzofuran

pg/g = picoograms per gram pg/L = picograms per liter RPD = relative percent difference

VQ = validation qualifier - See Attachment 2 for definitions

% = percent

a Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

^b No rejected data.

c A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

EVENT 1 FIELD DUPLICATE - DIOXIN PR1**DUP-01A

Positive Target Analyte Identification and Concentration Comparison^a

Analyte Identified	PR1WWDUP-01A Whole Water ^b (pg/L)	LQ°	VQ	PR1LDDUP-01A LSM Dissolved ^b (pg/L)	LQ°	VQ	PR1HDDUP-01A HSM Dissolved ^b (pg/L)	LQ°	VQ	% RPD	PR1LPDUP-01A LSM Particulate ^b (pg/g)	ιqʻ	VQ	PR1HPDUP-01A HSM Particulate ^b (pg/g)	LQ°	VQ	% RPD
2,3,7,8-TCDD														9.15		j	
1,2,3,7,8-PeCDD							The brain of				4.37	G		4.78	G	М	9
1,2,3,4,7,8-HxCDD	0.514	G					0.411	G			6.63	G		5.72		М	15
1,2,3,6,7,8-HxCDD	1.41	G		0.313	G		4.63			175	22.8	G		21.2		М	7
1,2,3,7,8,9-HxCDD	1.18	G					2.49	G			15.5	G		15.3		M	1
1,2,3,4,6,7,8-HpCDD	41.3		J	6.41		J	116		J	179	845		j	621		М	31
OCDD	429		J	44.0		J	720		J	177	8560		J	8960	D	М	5
2,3,7,8-TCDF							0.169	G			2.83	G		4.90		М	54
1,2,3,7,8-PeCDF				74			0.139	G			3.14	G		4.11	G	М	27
2,3,4,7,8-PeCDF	0.248	G					0.248	ø			8.37	G		5.26		М	46
1,2,3,4,7,8-HxCDF	1.33	G		0.364	G		0.912	G		86	13,3	G		31.5		J	81
1,2,3,6,7,8-HxCDF	1.29	G		0.373	G		1.04	G		94	14.0	G		18.2		M	26
2,3,4,6,7,8-HxCDF	1.39	G		0.321	G		0.922	G		97	15.8	G		20.9		М	28
1,2,3,7,8,9-HxCDF											4.55	G		1.89	G	M	83
1,2,3,4,6,7,8-HpCDF	20.5			3.20		j	17.6		J	138	215		J	271		M	23
1,2,3,4,7,8,9-HpCDF	1.56	G	J	0.363	G	J	1.45	G	J	120	16.2	G		18.7		M	14
OCDF	43.2		J	5.90		J	39.8		J	148	432		J	549		M	24

COPCs/COPECs listed in the FFS: 2,3,7,8-TCDD; 1,2,3,7,8-PeCDD; 1,2,3,6,7,8-HxCDD; 1,2,3,4,7,8-HxCDD; 1,2,3,7,8,9-HxCDD; 1,2,3,4,6,7,8-HpCDD; 0CDD; 2,3,7,8-TCDF; 1,2,3,7,8-PeCDF; 1,2,3,4,7,8-PeCDF; 1,2,3,4,7,8-HxCDF; 1,2,3,4,6,7,8-HxCDF; 1,2,3,4,7,8-HxCDF; 1,2,

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

FFS = focused fesability study HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

 ${\tt PCDD/PCDF = polychlorinated\,dibenzo-p-dioxin/polychlorinated\,dibenzofuran}$

pg/g = picoograms per gram

pg/L = picograms per liter

RPD = relative percent difference

VQ = validation qualifier - See Attachment 2 for definitions

% = percent

a Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

^b No rejected data.

^c A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Appendix B

Event #1, Attempt # 1 Results – PCB Congeners

Positive Target Analyte Identification and Concentration Comparison^a

	PR1CSOCLYWW-01A			PR1CSOCLYLD-01A LSM Dissolved ^b			PR1CSOCLYHD-01A HSM Dissolved ⁶				PR1CSOCLYLP-01A LSM Particulate ^b			PRICSOCLYHP-01A HSM Particulate ^b			
Analyte Identified	Whole Water ^b (pg/L)	LQ°	VQ	(pg/L)	ra	VQ	(pg/L)	ro,	VQ	% RPD	(pg/g)	rd,	VQ	(pg/g)	ra,	VQ	% RPD
PCB 1	26.3	D												156	D,G		
PCB 2														97.6	D,G		ž
PCB 3		D,G												182	D,G		
PCB 4/10	135	D				Substitution of the								870	D	М	
PCB 5/8		D												1340	D	M	
PCB 6	57.7	D		25.3	D,G									477	D,G	M	
PCB 7/9																	
PCB 11	422	D	j														
PCB 12/13																	
PCB 14																	
PCB 15	78.6	D												783	D	M	
PCB 16/32	222	B,D	J											2260	D	M	
PCB 17	121	B,D	J											1470	D	M	
PCB 18	296	B.D	J											2890	D	M	
PCB 19	63.9	D												568	D	M	
PCB 20/21/33	192	B,D	j											1130	D	J	
PCB 22	127	B,D	J											912	Б	Ü	
PCB 23	1	-,-	Ť													Ť ·	
PCB 24/27	31.1	D	 		-	+					524	D,G		315	D	М	49.8
PCB 25	52.0	D	 	13,3	D,G		43.6	Ь		106.5	916	D.		369	D	.1	85.1
PCB 26		D	+	13.3	0,0		40.0	U		100.5	1390	B,D		608	Б	11	78.3
	370	D	١,							_	6420			2620	D	0	
PCB 28	370	U	J		-	-		1			6420	B,D	J	2020	10	J	84.1
PCB 29			-													ļ	
PCB 30			1.					-									
PCB 31	309	B,D	J								6260	B,D	J	2280	D	J	93.2
PCB 34																<u> </u>	
PCB 35	27.0	D		6.22	D,G		16.4	D,G		90.0	474	B,D,G	3	197	D,G	J	82.6
PCB 36											145	D,G		75.1	D,G	J	63.5
PCB 37	110	B,D	J								1850	B,D	J	695	D	J	90.8
PCB 38																	
PCB 39																	
PCB 40	94.9	D									1610	D		835	D	M	63.4
PCB 41/64/71/72	449	B,D	j				215	B,D			8520	B,D	J	4210	D	M	67.7
PCB 42/59	157	D	j								2940	B,D	J	1350	D	М	74.1
PCB 43/49	415	B,D	J				195	B,D	J		7790	B,D	J	4070	D	M	62.7
PCB 44	568	B,D	J	300 00 00							10600	B,D	J	5490	D	M	63.5
PCB 45	79.3	D				100					1290	D		693	D	M	60.2
PCB 46	43.3	D									662	D		325	D	M	68.3
PCB 47	148	B,D	J														
PCB 48/75	75.1	B,D									1500	D		755	D	М	66.1
PCB 50		<u> </u>															
PCB 51	35.3	D												316	D	M	
PCB 52/69	822	B,D	.1				346	B,D	1		15500	B,D		8120	D	М	62.5
PCB 53	89.3	D	Ť	200				,,,			1550	D	-	736	D	M	71.2
PCB 54		Ť	 				2.16	D,G	1		,,,,			19.9	D,G		
PCB 55	15.2	D,G	1				2.10	0,0	~		232	D,G		175	D,G		28.0
PCB 56/60	340	B,D	1.1				129	B,D			6910	B.D		3180	D,G	M	73.9
PCB 57	J40	0,0	 				12.0	0,0			00.10	טיי	-	51.6	D,G	a construction	13.3
	_	-	+							 				01.0	۵,۷	ivi	
PCB 58	017	B,D	1				200	0.0		 	15700	B,D		9290	h	8.4	1 50.5
PCB 61/70	817	ח,ם	J				309	B,D			10700	0,0	J	8380	D	M	60.8
PCB 62	20.4	<u> </u>	-				0.70	h			470			070	1		⊢
PCB 63	23.1	D					9.78	D,G			476	D,G		270	D	M	55.2
PCB 65																	
PCB 67		D,G									271	D,G		134	D,G		67.7
PCB 68	4.85	B,D,G												49.4	D,G	M	
PCB 73																	
PCB 74	242	B,D	J	44.2	D		102	D		79.1	4730	B,D	J	2360	D	M	66.9
PCB 76/66	552	B,D	J				207	D			10700	B,D	J	5110	D	M	70.7

	1		1	PR1CSOCLYLD-01A			PR1CSOCLYHD-01/				PR1CSOCLYLP-01A	1		PRICSOCLYHP-01A	ı		
	PR1CSOCLYWW-01A			LSM Dissolved ^b			HSM Dissolved ^b	`			LSM Particulate ^b			HSM Particulate ^b			
Analyte Identified	Whole Water ^b (pg/L)	LQ ^c	VQ	(pg/L)	ro.	VQ	(pg/L)	LQ	VQ	% RPD	(pg/g)	ra	VQ	(pg/g)	LQ°	VQ	% RPD
PCB 77	72.3	B,D	1											924	D	М	
PCB 78																	
PCB 79	16.3	B,D,G		5.38	D,G		7.18	D,G		28.7				251	D	M	
PCB 80																	
PCB 81	12.5	D,G												95.7	D,G	M	
PCB 82	228	D	J				58.3	D			4460	D	J	2890	D	J	42.7
PCB 83			ļ														
PCB 84/92	674	B,D	J				197	B,D	J		12600	D	J	8330	D	J	40.8
PCB 85/116	215	D	J				47.6	D	J		4210	D	J	2690	D	J	44.1
PCB 86 PCB 87/117/125	677	D D	J				193	D			11500	D		8010	D		25.0
PCB 88/91	209	B,D D	J				54.2	ם	J		3680	ם	J	2330	D	J	35.8 44.9
PCB 89	18.5	D,G	1				34.2	U	J		327	D,G	1	174	D,G	J	61.1
PCB 90/101	1660	B,D	.1		 	 	466	B,D	1		30700	B,D	1	20200	D	i i	41.3
PCB 93	1000	0,0	1		-		100	+10,0			00,00	10,5	-	LULUU		-	1
PCB 94																	
PCB 95/98/102	1180	B,D	J								20900	B,D	J	14000	D	J	39.5
PCB 96		<u> </u>									181	D,G	J	128	D,G	J	34.3
PCB 97	520	D	j				156	D	J		9390	D	J	6330	D	J	38.9
PCB 99	607	B,D	j				177	D	J		11200	D	J	7960	D	J	33.8
PCB 100				200													
PCB 103											124	D,G	J	124	D,G	J	0.0
PCB 104																	
PCB 105	684	D	J				177	D			11600	B,D	J	8250	D	J	33.8
PCB 106/118	1560	B,D	J				401	B,D			29000	B,D	J	20100	D	J	36.3
PCB 107/109	74.7	D	j				22.3	D			1750	D	J	1100	D	J	45.6
PCB 108/112	72.6	D	J	14.6	D,G		22.7	D	J	43.4	1260	D	J	935	D	J	29.6
PCB 110	1670	B,D	J				423	B,D	J		31200	B,D	J	20000	D	J	43.8
PCB 111/115	23.0	D	J								650	D	J	286	D	J	77.8
PCB 113			ļ. —														
PCB 114	34.1	D	J			100	12.1	D,G	J		658	D	J	557	D	J	16.6
PCB 119	22.4	D	J				6.77	D,G	J		429	D,G	J	263	D	J	48.0
PCB 120 PCB 121			-									-			-		
PCB 122	15.5	D	1					+			402	D,G	1	221	D,G		58.1
PCB 123	13.3	 	1							-	628	D,G	1	301	D.G	J	70.4
PCB 124	73.7	D	i.i			100	19.7	D,G			1450	D	ı	960	Б	.1	40.7
PCB 126	20.7	D,G					10.1	10,0			1400	1	•	261	D	J	40.7
PCB 127	20.7	5,0	ľ												Ť		
PCB 128/162	376	B,D	J				82.2	Ь						5210	D	J	
PCB 129	129	B,D	J				33.6	D						1740	D	J	
PCB 130	117	B,D	J				25.7	D						1720	D	J	
PCB 131																	
PCB 132/161	532	B,D	J				122	D			9660	D	J	7190	D	J	29.3
PCB 133/142	61.6	D	J				15.3	D,G			962	D	J	748	D	J	25.0
PCB 134/143	120	D	J				29.8	D			1880	D	J	1480	D	J	23.8
PCB 135	156	D	J				48.2	D						2020	D	J	
PCB 136	169	D	J				41.9	D						1880	D	J	
PCB 137	73.7	D	J				18.5	D,G				<u> </u>		854	D	J	
PCB 138/163/164	1990	B,D	J				426	B,D			32800	B,D	J	25100	D	ل	26.6
PCB 139/149	1040	D	J				300	B,D			19700	D	J	13700	D	J	35.9
PCB 140	050	D D			9000	<u> </u>	00.0	-			00.40			10.10			
PCB 141	358	B,D	J				83.8	D			6340	D		4640	D	J	31.0
PCB 144 PCB 145	57.2	D	J		-		18.4	D,G		-	1170	D	J	873	D	J	29.1
PCB 145 PCB 146/165	200	<u> </u>	J				50.2	D		—	3510	D		2500	D		22.6
PCB 147	200 22.8	D D	i i				50.2	U		—	3510 420		J	273	D	J	33.6 42.4
PCB 148	22.0	۲	۲							-	740	0,6	J	213	10	J	42.4
PCB 150	+	 	1														
PCB 151	255	D	J		 						4450	D	J	2960	D	i.	40.2
PCB 152	-	Ť	Ť										-		Ē		1.5.2
PCB 153	1440	B,D	J				360	B,D			24100	B,D	J	16700	D	J	36.3
		D,G	i.	\$10002174 D-01218 BROOKS BROOKS	- Constitution of the Cons	A LANGUAGE CONTRACTOR OF THE PARTY OF THE PA		NAME OF TAXABLE PARTY OF TAXABLE PARTY.	Programme College		178	D,G	S. W. Williams	146	D,G	10000000	19.8

				PR1CSOCLYLD-01A			PR1CSOCLYHD-01A				PR1CSOCLYLP-01A			PRICSOCLYHP-01A			
Analyte Identified	PR1CSOCLYWW-01A Whole Water ^b (pg/L)	LQ°	VQ	LSM Dissolved ^b (pg/L)	LQ'	VQ	HSM Dissolved ^b (pg/L)	LQ°	VQ	% RPD	LSM Particulate ^b (pg/g)	ια	VQ	HSM Particulate ^b (pg/g)	LQ'	VQ	% RPD
PCB 155		D,G	J									D.G		148	D.G	J	53.1
PCB 156	218	B.D	j				46.7	D						3140	D	J	
PCB 157	56.6	B.D	J	0.000										758	n	j d	
PCB 158/160	243	B.D	J				48.3	D						2950	ā	ī	
PCB 159	2.10	0,0	 				10.0	-							1		
PCB 166			1											181	D.G		
PCB 167	95.1	B,D	1.1				20.5	D.G						1360	D	ñ	
PCB 168	00.1	0,0	 			1	20.0	0,0						1000	۳-	-	
PCB 169			+		-					_					-		
PCB 170	365	B,D	1				92.0	D			6300	D	1	5570	D	h —	12.3
PCB 171	102	D	1.1				23.5	D			1740	D	1	1420	Б	1	20.3
PCB 172	61.7	D	J				23.3	U				D	1	833	D	-	13.3
PCB 173	01.7	<u> </u>	10								932	<u> </u>	J	000	10	J	15.5
PCB 174	413	D	1				94.3	D		-	6990	D		5140	D	1	20.5
PCB 175	413	D	J			+	94.3	U		-	262	D.G	J	186	4	J 1	30.5
	44.0	D	1,	7.07	50		44.0	0.0		20.0		INCOME TO SECU	J		D,G	-	33.9
PCB 176		_	J	7.87	D,G		11.6	D,G		38.3	731	D	0.0000000000000000000000000000000000000	608	D	J.	18.4
PCB 177		B,D	J				55.8	D			4110	D	J	3180	D	J	25.5
PCB 178	70.8	D	J		<u> </u>						1020	D	ل	877	D	J	15.1
PCB 179	165	D	J								2450	D	J	2030	D	J	18.8
PCB 180	889	B,D	j								15200	D	J	11400	D	J	28.6
PCB 181																	
PCB 182/187	388	B,D	J								6190	D	J	4870	D	J	23.9
PCB 183	177	D	J								2990	D	J	2260	D	J	27.8
PCB 184	18.5	D,G	J	0.00							299	D,G	J	217	D,G	J	31.8
PCB 185	43.8	D	j				11.1	D,G			694	D	J	519	D	J	28.9
PCB 186																	
PCB 188																	
PCB 189																	
PCB 190	99.0	D	j	8.57	D,G		16.8	D,G		64.9	1220	D	J	1060	D	J	14.0
PCB 191														217	D,G	J	
PCB 192																	
PCB 193	66.0	D	J								685	D	J	510	D	J	29.3
PCB 194	191	D	j								3110	B,D	J	2540	D	J	20.2
PCB 195	86.2	D	J								1160	D	J	1310	D	J	12.1
PCB 196/203	152	D	J								3260	D	J	1900	D	J	52.7
PCB 197														112	D,G	J	
PCB 198																	
PCB 199	165	B,D	1.1								2730	D		2060	D	1	28.0
PCB 200	24.6	D	i.								426	D.G	1				
PCB 201	29.3	D	.1		1						462	D.G	ì	353	Б		26.7
PCB 202	44.8	D	1.1	4.88	D.G					 	813	D.	1	561	Б	fi d	36.7
PCB 204	7 1.0	Ε	Ť		-										1		———
PCB 205			+														\vdash
PCB 206	132	B.D	+							 	2680	D	1	1930	D		32.5
PCB 207	102	0,0	+							—	253	D.G	1	166	D.G	-	41.5
PCB 207	49.0	D	+-							_	1000	D,G	-	608	D,G	1	
			J i			-					1000	U	J			9	48.8
PCB 209	94.7	B,D	J											1410	D	J	

COPCs/COPECs listed in the FFS: PCB -77, PCB -81, PCB -105, PCB -114, PCB -118, PCB -123, PCB -126, PCB -156, PCB -157, PCB -167, PCB -169, and PCB -189.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study
HSM = high-solids mass

LSM = low-solids mass $LQ = laboratory \ qualifier \cdot See \ Attachment \ 1 \ for \ definitions \\ PCB = polychlorinated \ biphenyl \\ pg/g = picograms \ per \ gram \\$

pg/L = picograms per liter RPD = relative percent difference VQ = validation qualifier - See Attachment 2 for definitions % = percent

a Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

^b No rejected data.

^c A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Positive Target Analyte Identification and Concentration Comparison^a

	PR1WWDUP-02B Whole Water ^b			PR1LDDUP-02B			PR1HDDUP-02B				PR1LPDUP-02B			PR1HPDUP-02B			1
Analyte Identified	(pg/L)	IQ°	VQ	LSM Dissolved b (pg/L)	LQ*	VQ	HSM Dissolved ^b (pg/L)	LQ°	VQ	% RPD	LSM Particulate ^b (pg/g)	ro,	VQ	HSM Particulate ^b (pg/g)	LQ°	VQ	% RPD
· · · · · · · · · · · · · · · · · · ·				(bg/r)	- cu	l vu	(1987-1)	1 LQ	VQ	70 KPD	(1478)	LU	VQ			99619,010821	70 KPD
PCB 1 PCB 2	24.6	D	J			-		-						158 93.7	D,G D,G	M	⊢—
PCB 3	15.3	D,G						-						193		M M	!
PCB 4/10	103	D,G	J	70										804	D,G	M	
PCB 5/8	104	D	J			-								1270	D	M	
PCB 6	36.8	D,G	J	23.6	D,G	1,		1						438	A-27,002	M	-
PCB 7/9	30.0	D,G	3	23.0	D,G	U	+	-						430	۵,0	IVI	_
PCB 11	280	D	J					-								-	.
PCB 12/13	200	10	3		100		12.1	D,G									
PCB 14		 					14.1	0,0									
PCB 15	30.6	Ь	J				-	-						706	D	м	
PCB 16/32	160	B,D	J					+						2180	D	M	—
PCB 17	78.2	B,D	J			-								1400	D	M	
PCB 18	180	B,D	J			-		-						2830	D	M	
	49.0	D D	1		-			1						581	D	M	
PCB 19 PCB 20/21/33	104	B,D	j			1		1	-					1050	D	IVI I	⊢—
PCB 20/21/33 PCB 22	81.2	B,D	J			+								679	D.	J J	—
PCB 23	01.2	ID,U	U											019	U	J	-
	14.2	D.G.	+					-		-				305	D	8.4	—
PCB 24/27 PCB 25	14.2 34.1	D,G D	+	15.8	D.G.	-	44.7	D		95.5	293	D,G		305 344	ם ח	M	16.0
PCB 25	46.3	D	1	10.0	D,G	6,0,000	44.1	ח		95.5	೭೪೦	<u>ا</u> ي.ن		344 446	D	J	16.0
		D													075233023000	J	-
PCB 28	217	D	J											2880	D	J	
PCB 29		-						1									-
PCB 30	242	-	1.					1						2000			-
PCB 31	210	B,D	J		-			-						2260	D	J	—
PCB 34	45.0						10.4	-							_		
PCB 35	15.0	D,G					12.4	D,G						244	D	J	—
PCB 36	50.0	-	1.	300,000				1						63.2	D,G D	J	—
PCB 37	59.9	B,D	J					-						861	100000000000000000000000000000000000000	J	—
PCB 38														96.4	D,G	J	⊢—
PCB 39	50.4					-								700			-
PCB 40	56.1	D				-	007	0.0			0000			769	D	M	
PCB 41/64/71/72	251	B,D	J				207	B,D	J		2680	B,D	J	3810	D	M	34.8
PCB 42/59	74.8	D	J								906	B,D	J	1260	D	M	32.7
PCB 43/49	224	B,D	J				203	B,D	J I		2500	B,D	J	3640	D	J	37.1
PCB 44	234	B,D	J			1	251	B,D	J		3440	B,D	J	4830	D	M	33.6
PCB 45	45.5	D						1						557	D	J	
PCB 46	20.0	D,G									196	D,G		301	D	J	42.3
PCB 47	85.7	B,D	J											004			Ь——
PCB 48/75	45.7	B,D	1			+		-	-					694	D	М	—
PCB 50	20.2	ln.c	1				9.90							244	D	1	—
PCB 51	20.2	D,G	.1				200	l			E440			244	Company of the last	J	
PCB 52/69	459	B,D	J			-	362	B,D	J		5110	B,D	J	7500	D	J	37.9
PCB 53	46.5	D	1			+	53.7	D	J					658	D	J	—
PCB 54	10.6	l	1							ļ	05.0	0.0		400	0.0		
PCB 55 PCB 56/60	10.6	D,G B,D	1.1			-	128	D.C.			85.3 2050	D,G B,D		186 3160	D,G D	M	74.2
	188	ח'מ	J				128	B,D		-	2030	ט,ט	J	3 100	U	М	42.6
PCB 57	_	1	+							-							200.0
PCB 58	446	<u> </u>	1				206	100			4020	0.0		7040	D		
PCB 61/70	446	B,D	J			-	296	B,D			4930	B,D	J	7940	D	М	46.8
PCB 62	10.0	10.0	1					1		ļ	4.47						L
PCB 63	12.2	D,G					12.3	D,G			147	D,G		214	D,G	M	37.1
PCB 65	7.70	I	1					-			54.4						— —
PCB 67	7.72	D,G	1			1		-			81.1	D,G		115	D,G	M .	34.6
PCB 68		1	1					-									Ь——
PCB 73			1														Ь—
PCB 74	137	B,D	J	46.3	D	J	96.8	D		70.6	1500	B,D	J	2180	D	M	37.0

	PR1WWDUP-02B			PR1LDDUP-02B			PR1HDDUP-02B				PR1LPDUP-02B			PR1HPDUP-02B			
	Whole Water ^b			LSM Dissolved b			HSM Dissolved ^b				LSM Particulate ^b			HSM Particulate ^b			1 1
Analyte Identified	(pg/L)	LQ°	VQ	(pg/L)	LQʻ	VQ	(pg/L)	LQʻ	VQ	% RPD	(pg/g)	ΙQʻ	VQ	(pg/g)	LQ°	VQ	% RPD
PCB 76/66	302	B,D	J				213	D			3220	B,D	J	5000	D	M	43.3
PCB 77	44.7	B,D												1010	D	М	
PCB 78																	ldot
PCB 79	11.7	B,D,G												253	D	M	lacksquare
PCB 80															<u> </u>		ldot
PCB 81	7.36	D,G												113	D,G	M	lacksquare
PCB 82	107	D	J				70.6	D				<u> </u>		2690	D	J	igsquare
PCB 83												<u> </u>					
PCB 84/92	380	B,D	J				208	B,D			-222	<u> </u>		8250	D	J	
PCB 85/116	92.8	D	J				58.6	D			1360	D	J	2560	D	J	61.2
PCB 86			1.					<u> </u>							-		lacksquare
PCB 87/117/125	388	B,D	J				195	D			3710	D	J	7820	D	J	71.3
PCB 88/91	108	D	J				61.1	D	J		1210	D	J	2190	D	J	57.6
PCB 89	9.48	D,G					100	-			10000	<u> </u>		00400	_		\vdash
PCB 90/101	920	B,D	J		ļ		488	B,D			10300	B,D	J	20100	D	J	\vdash
PCB 93								-							ļ	4	lacksquare
PCB 94	077	D D		100							7050	<u> </u>		10000	<u> </u>		L
PCB 95/98/102	677	B,D	J								7250	B,D	J	12300	D	J	51.7
PCB 96		_	1,											118	D,G	J	-
PCB 97	299	D	J		600000000000000000000000000000000000000		151	D			0000	_		6100	D	J	
PCB 99	341	B,D	J				185	D			3690	D	J	7950	D	J	73.2
PCB 100							4.00	<u> </u>	,						-		
PCB 103							4.69	D,G	J						-		
PCB 104	255	_					400	In.				ļ		0400	<u> </u>	ļ. —	
PCB 105	355	D B,D	J				182	D			0400	0.0		8120 21000	D D	.1	H
PCB 106/118 PCB 107/109	821	D D	J	45.0	D 0		405	B,D			9100	B,D	J J		D D	J J	79.1
PCB 107/109 PCB 108/112	40.7 41.0	D		15.8 14.9	D,G D,G		24.0	h		50.2	592 441	D D	J	1020 893	ם מו	J	53.1
	859		J	14.9	D,G		24.9	D		50.3	441	η		19900		J	67.8
PCB 110 PCB 111/115	18.9	B,D D,G	J				457	B,D			241	D,G		314	D D	J .	25.2
PCB 1117113	10.9	D,G	J	0.000	-			+			241	0,6		314	U	J	26.3
PCB 114	20.4	D,G	J								208	D,G		459	D	1	75.2
PCB 114 PCB 119	14.8	D,G	J								139	D,G		323	D D	J	75.3 79.7
PCB 120	14.0	D,G	3					-			139	D,G		020	Ρ	-	/3./
PCB 121												-					-
PCB 122											114	D,G			 		\vdash
PCB 123											189	D,G		322	D	i.i	52.1
PCB 124	37.1	D		12.5	D,G						446	D		969	D D		73.9
PCB 126	10.3	D,G		TE.S	0,0						7,10			278	D	i	15.5
PCB 127	10.0	-,-													 		
PCB 128/162	184	B,D	J				81.6	D						5050	D	J	
PCB 129	67.1	B,D	J				27.8	D						1670	D	J	
PCB 130	48.8	B,D	J											1600	D	J	
PCB 131		l															\Box
PCB 132/161	274	B,D	J				128	D	7,02					7060	D	J	\Box
PCB 133/142	28.6	D					14.8	D,G						775	D	J	\Box
PCB 134/143	64.2	D	J				32.1	D						1540	D	J	\Box
PCB 135	96.2	D	J				43.5	D						1990	D	J	
PCB 136	84.3	D	J				40.0	D						1990	D	J	
PCB 137	37.1	D												1300	D	J	
PCB 138/163/164	922	B,D	J				426	B,D						24300	D	J	
PCB 139/149	601	D	J			70 E 180	249	B,D						13200	D	J	
PCB 140																	
PCB 141	170	B,D	J				83.0	D						4540	D	J	
PCB 144	29.9	D					16.6	D,G	10		349	D		676	D	J	
PCB 145																	
PCB 146/165	102	D	J				51.7	D						2530	D	J	
PCB 147							6.66	D,G			199	D,G		297	D	J	
PCB 148						10.0											
PCB 150					15												
PCB 151	138	D	J								1440	D	J	3120	D	J	73.7

	PR1WWDUP-02B			PR1LDDUP-028			PR1HDDUP-02B				PR1LPDUP-02B			PR1HPDUP-02B			
Analyte Identified	Whole Water ^b (pg/L)	LQ ^c	VQ	LSM Dissolved b (pg/L)	£Q°	vq	HSM Dissolved ^b (pg/L)	LQ°	VQ	% RPD	LSM Particulate ^b (pg/g)	ια·	VQ	HSM Particulate ^b (pg/g)	ιqʻ	VQ	% RPD
PCB 152																	
PCB 153	690	B,D	J				346	B,D						18200	D	J	
PCB 154		-,-									97.4	D,G		142	D,G	Ĵ	37.3
PCB 155	8.66	D,G					10.5	D,G									
PCB 156	106	B,D	J				44.1	D						3050	D	J	
PCB 157	22.3	B,D												711	D	J	
PCB 158/160	118	B,D	J				53.5	D						3050	D	J	
PCB 159																	
PCB 166						1000											
PCB 167	39.2	B,D	J				20.3	D,G						1300	D	J	
PCB 168																	
PCB 169					- 200												
PCB 170	162	B,D	J				101	D						5170	D	J	
PCB 171	48.0	D	J											1360	D	J	
PCB 172	25.5	D												773	D	J	
PCB 173														143	D,G	J	
PCB 174	181	D	J			1000	102	D						4970	D	J	
PCB 175					1000									216	D,G	J	
PCB 176	18.6	D,G		7.51	D,G	100000	10.3	D,G		31.3	202	D,G		547	D	J	92.1
PCB 177	108	B,D	J				48.9	D						3020	D	J	
PCB 178	25.4	D	J											936	D	J	
PCB 179	73.8	D	J								685	D	J	1920	D	J	94.8
PCB 180	396	B,D	J											11500	D	J	
PCB 181																	
PCB 182/187	163	B,D	J											5030	D	J	and the second
PCB 183	79.1	D	J											2290	D	J	
PCB 184	13.6	D,G		7.16	D,G		16.5	D,G		79.0	124	D,G		209	D,G	J	51.1
PCB 185	18.9	D,G					13.7	D,G			207	D,G		532	D	J	88.0
PCB 186							1.00										
PCB 188																	
PCB 189														251	D	J	
PCB 190	37.7	D	J	12.3	D,G		19.7	D,G		46.3	378	D		1010	D	J	91.1
PCB 191											93.0	D,G		172	D,G	J	59.6
PCB 192																	
PCB 193	21.0	D	J								198	D,G		484	D	J	83.9
PCB 194	80.4	D	J								906	B,D	J	2420	D	J	91.0
PCB 195	29.4	D	J	10.2	D,G	J					342	D	J	1050	D	J	101.7
PCB 196/203	69.5	D	J											2080	D	J	
PCB 197									3								of the same
PCB 198	7.75	D,G															
PCB 199	87.0	B,D	J								736	D	J	2110	D	J	96.6
PCB 200	9.54	D,G									112	D,G		292	D	J	89.1
PCB 201											110	D,G	Z	287	D	J	89.2
PCB 202	18.6	D,G									193	D,G		587	D	J	101.0
PCB 204																	
PCB 205																	
PCB 206	61.0	B,D	J											2110	D	J	
PCB 207											77.0	D,G	J				
PCB 208	17.7	D,G									192	D,G	J	621	D	J	105.5
PCB 209	29.6	B,D,G	J											1380	D	J	

COPCs/COPECs listed in the FFS: PCB -77, PCB -81, PCB -105, PCB -114, PCB -118, PCB -123, PCB -126, PCB -156, PCB -157, PCB -167, PCB -169, and PCB -189.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern FFS = focused fesability study HSM = high-solids mass LSM = low-solids mass LQ = laboratory qualifier - See Attachment 1 for definitions PCB = polychlorinated biphenyl pg/g = picograms per gram

pg/L = picograms per liter
RPD = relative percent difference
VQ = validation qualifier - See Attachment 2 for definitions
% = percent

a Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

^b No rejected data.

^c A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier results fall below the low point of the calibration curve.

Appendix C Contingency Samples Used During CSO Phase I Sampling Events

Contingency Samples Used During the CSO Phase I Sampling Events

				Reason for Cor			
SDG#	Contingency Sample Bottles Used	Sample Type	Lab Received Broken	Lab Received Outside Temperature	Lost in transit to Lab	Re- analysis Required	Notes
PR105	1	WW PCB Congener			NA	NA	Event 1 Attempt 1
PR105	32	WW Dioxin/Furan			NA		Event 1 Attempt 1
PR107	0	HSM Dissolved Dioxin/Furan	NA	NA	NA	NA	Dioxin analysis 1613B- 1 contingency sample bottle was received broken. No contingency bottle was used in extraction. Event 1 Attempt 1
PR107	2	HSM Dissolved PCB Congener		NA	NA	NA	Event 1 Attempt 1
PR134	4	WW Pesticide	NA	NA	**************************************	NA	Event 2 Attempt 2
PR134	16	WW Dioxin/Furan	NA	NA		NA	Event 2 Attempt 2
PR134	8	WW PCB Congener	NA	NA		NA	Event 2 Attempt 2
PR145	4	WW Dioxin/Furan		NA	NA	NA	Event 1 Attempt 3

Notes:

HSM= High Solids Mass

NA= Not Appicable

SDG = Sample Delivery Group

WW = Whole Water

^		pe		ᆚ	:	
м	E 31	и.	11	"	ΙX	

CSO/SWO Phase I Field Blank Contamination Results

CSO/SWO Phase I Field	d Blank Contaminat	ion Results Quali	fied
	Number of Samples Affected	Number of Results Affected	Percent of the Total Results Affected
HSM Particulate			
Semivolatiles	1	2	1.0
Organochlorine Pesticides	4	20	17.9
Semivolatiles SIM	2	8	6.7
Cyanide	3	3	75.0
PCDD/PCDFs	3	5	4.9
PCB Congeners	3	22	2.2
Chlorinated Herbicide	6	10	42.0
HSM Dissolved			
Semivolatiles	3	4	2.0
Organochlorine Pesticides	4	32	28.6
Semivolatiles SIM	4	35	29.2
PCDD/PCDFs	2	9	8.8
PCB Congeners	6	305	30.3
Chlorinated Herbicide	2	7	29.2
TOC	2	2	50.0
TEPH	2	2	50.0
TSS	2	2	25.0
TDS	2	2	25.0
LSM Particulate	_		23.0
Semivolatiles	3	5	2.5
Organochlorine Pesticides	4	33	29.5
Semivolatiles SIM	4	28	23.3
PCDD/PCDFs	3	8	7.8
PCB Congeners	6	275	27.3
LSM Dissolved	U	2/3	27.5
	2		2.0
Semivolatiles	3	4	2.0
Organochlorine Pesticides	4	30	26.8
Semivolatiles SIM	4	26	21.7
PCDD/PCDFs	4	10	9.8
PCB Congeners	6	366	36.3
Chlorinated Herbicide	4	9	37.5
DOC	4	4	100.0
Whole Water			T
Semivolatiles	4	4	2.0
Organochlorine Pesticide	4	29	25.9
Semivolatiles SIM	3	23	19.2
Metals	4	6	6.5
Cyanide	2	2	50.0
PCDD/PCDFs	2	7	6.9
PCB Congeners	5	123	12.2
Chlorinated Herbicide	4	7	29.2
TOC	2	2	50.0
TDS	2	2	50.0
Grab Water Dissolved			
Metals	4	8	8.7

Notes:

CSO/SWO = combined sewer overflow/stormwater outfall

DOC = dissolved organic carbon

 $\mathsf{HSM} = \mathsf{high}\text{-}\mathsf{solids}$ mass

LSM = low-solids mass

PCB = polychlorinated biphenyl

SIM = selective ion monitoring

TDS = total dissolved solids

TEPH = total extractable petroleum hydrocarbons

TOC = total organic carbon

TSS = total suspended solids

Appendix E

Field Blank Results Assessment

1. Polychlorinated Dibenzo-p-dioxins/Polychlorinated Dibenzofurans

Table 1
PCDD/PCDF¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection
Method for Event #2, Attempt #2

	PR10	CSOCLY	**-02B	PI	R1**DUP-	02B
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³
Detections Reported by Laboratory	10	8	15	12	14	16
Detections Impacted by Field Blank ⁴	3	4	1	4	3	1
Usable Results ⁵	7	4	14	8	11	15

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-25, Rev. 3, 2006
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- Based on Worksheet #11

The final recommended PCDD/PCDF sample collection method (HSM) was not impacted by the field blank concentrations during the Event #2, Attempt #2 for either the primary or duplicate sample.

Table 2
PCDD/PCDF¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection
Method for Event #1, Attempt #3

	PR10	CSOCLY	**-01C	PI	R1**DUP-	01C
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³
Detections Reported by Laboratory	14	15	16	13	15	15
Detections Impacted by Field Blank ⁴	0	0	1	0	0	0
Usable Results ⁵	14	15	15	13	15	15

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-25, Rev. 2, 2006
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

The final recommended PCDD/PCDF sample collection method was not impacted by the field blank concentrations during the Event #1, Attempt #3 primary sample (inconclusive) or duplicate sample (LSM/HSM).

2. Polychlorinated Biphenyl Congeners

Table 3
PCB Congeners¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection
Method for Event #2, Attempt #2

	PR10	CSOCLY	**-02B	PI	R1**DUP-	02B
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³
Detections Reported by Laboratory	6	11	9	8	9	10
Detections Impacted by Field Blank ⁴	0	4	0	1	1	1
Usable Results ⁵	6	7	9	7	8	9

Notes:

- 1. Validation Guidance-EDS SOP: Congener PCB, Rev. 3, July 2010
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- Based on Worksheet #11

The final recommended PCB congener sample collection method (HSM) was not impacted by the field blank concentrations during the Event #2, Attempt #2 for the duplicate sample. The number of positive COPCs/COPECs reported (as well as overall target analytes detected) is significantly higher in the HSM duplicate sample than the other sample collection methods with and without qualification for associated field blank concentrations. However, the field blank detections associated with the primary sample impacted the final recommended sample collection method (HSM) as indicated in the table above.

Table 4
PCB Congeners¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection
Method for Event #1, Attempt #3

	PR1	CSOLLY	**-01C	PF	11**DUP-01C		
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³	
Detections Reported by Laboratory	7	8	9	7	6	9	
Detections Impacted by Field Blank ⁴	1	0	0	1	1	0	
Usable Results ⁵	6	8	9	6	5	9	

Notes:

- 1. Validation Guidance-EDS SOP: Congener PCB, Rev. 3, July 2010
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

The final PCB congener recommended sample collection method (HSM) was not impacted by the field blank concentrations during the Event #1, Attempt #3 in either the primary or duplicate sample.

3. Aroclor Polychlorinated Biphenyls

No field blank concentrations were present. Field blank results did not impact any positive result reported during Phase I for the Aroclor PCBs for LSM, HSM, or whole water collection methods.

4. Organochlorine Pesticides

Table 5
Organochlorine Pesticides¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection Method for Event #1, Attempt #2

	PR10	CSOCLY	**-01B	PF	R1**DUP-	01B
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³
Detections Reported by Laboratory	6	6	N/A	6	6	6
Detections Impacted by Field Blank ⁴	3	3	1	3	3	1
Usable Results ⁵	3	3	N/A	3	3	5

Notes:

- 1. Validation Guidance EDS SOP: Organochlorine Pesticides by HRGC/HRMS USEPA 1699, Rev. 0 7/10
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

N/A = not applicable; sample was eliminated from further evaluation during Step 2, because less than 90% usable data were obtained

The final organochlorine pesticide recommended sample collection method (inconclusive) was not impacted by the field blank concentrations during Event #1, Attempt #2 in the primary sample. However, the field blank concentrations associated with the duplicate sample for COPCs/COPECs impacted the final recommended sample collection method (HSM) as indicated in the table above.

Table 6
Organochlorine Pesticides¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection Method for Event #2, Attempt #2

	PR1CSOCLY**-02B			PF	R1**DUP-02B			
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³		
Detections Reported by Laboratory	6	6	6	6	6	6		
Detections Impacted by Field Blank ⁴	3	3	2	3	3	3		
Usable Results ⁵	3	3	4	3	3	3		

Notes:

- Validation Guidance EDS SOP: Organochlorine Pesticides by HRGC/HRMS USEPA 1699, Rev. 0 7/10
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

The final recommended organochlorine pesticide sample collection method (inconclusive) was not impacted by the field blank concentrations during Event #2, Attempt #2 for the duplicate sample. However, the field blank concentrations associated with the primary sample for COPCs/COPECs impacted the final recommended sample collection method (HSM) as indicated in the table above.

5. Semivolatile Organic Compounds

There are no COPC/COPECs in the target list for SVOCs. Therefore, the following tables compare the analytes affected by the field blank results with the Target Analyte List (TAL).

Table 7
SVOCs¹ – Target Analytes Impacted by Field Blank Concentrations by Collection Method for Event #1, Attempt #2

	PR1CSOCLY**-01B			PF	R1**DUP-01B		
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³	
Detections Reported by Laboratory	5	N/A	N/A	5	5	N/A	
Detections Impacted by Field Blank ⁴	1	2	0	1	1	0	
Usable Results ⁵	4	N/A	N/A	4	4	N/A	

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-35, Rev.1, August, 2007
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

N/A = not applicable; sample was eliminated from further evaluation during Step 2, because less than 90% usable data were obtained

The final recommended SVOC sample collection method was not impacted by the field blank concentrations during Event #1, Attempt #2 for either the primary (inconclusive) or duplicate sample (inconclusive).

Table 8
SVOCs¹ – Target Analytes Impacted by Field Blank Concentrations by Collection Method for Event #2, Attempt #2

	PR1CSOCLY**-02B			PI	R1**DUP-02B			
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³		
Detections Reported by Laboratory	5	6	10	5	6	8		
Detections Impacted by Field Blank ⁴	1	1	0	1	1	0		
Usable Results ⁵	4	5	10	4	5	8		

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-35, Rev.1, August, 2007
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

The final recommended SVOC sample collection method was not impacted by the field blank concentrations during Event #2, Attempt #2 for either the primary (HSM) or duplicate sample (inconclusive).

6. Semivolatile Organic Compounds Selective Ion Monitoring

Table 9
SVOCs SIM¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection
Method for Event #1, Attempt #2

	PR1CSOCLY**-01B			PI	R1**DUP-01B			
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³		
Detections Reported by Laboratory	16	17	16	16	16	15		
Detections Impacted by Field Blank ⁴	4	7	0	7	5	1		
Usable Results ⁵	12	10	16	9	11	14		

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-35, Rev.1, August, 2007
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- Based on Worksheet #11

The final recommended SVOC SIM sample collection method was impacted by the field blank concentrations during Event #1, Attempt #2 for both the primary (HSM) and duplicate samples (HSM).

Table 10
SVOC SIM¹ – COPCs/COPECs Analytes Impacted by Field Blank Concentrations by Collection Method for Event #2, Attempt #2

	PR1CSOCLY**-02B			PF	R1**DUP-02B		
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³	
Detections Reported by Laboratory	16	17	17	17	17	17	
Detections Impacted by Field Blank ⁴	1	1	0	0	1	0	
Usable Results ⁵	15	16	17	17	16	17	

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-35, Rev.1, August, 2007
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- Based on Worksheet #11

The final recommended SVOC SIM sample collection method was not impacted by field blank concentrations during Event #2, Attempt #2 for either the primary (inconclusive) or duplicate samples (inconclusive).

7. Chlorinated Herbicides

There are no COPC/COPECs in the target list for chlorinated herbicides. Therefore, the following tables compare the analytes affected by the field blank results with the chlorinated herbicide TAL.

Table 11
Chlorinated Herbicides¹ – Target Analytes Impacted by Field Blank Concentrations by Collection
Method for Event #1, Attempt #2

	PR1CSOCLY**-01B			PI	R1**DUP-01B			
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³		
Detections Reported by Laboratory	1	4	3	0	3	3		
Detections Impacted by Field Blank ⁴	1	2	2	0	2	2		
Usable Results ⁵	0	2	1	0	1	1		

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-17, Rev.3, July, 2008
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

The final recommended chlorinated herbicide sample collection method was not impacted by field blank concentrations during Event #1, Attempt #2 for either the primary (LSM) or duplicate samples (LSM/HSM).

Table 12

Chlorinated Herbicides¹ – Target Analytes Impacted by Field Blank Concentrations by Collection Method for Event #2, Attempt #2

	PR1CSOCLY**-02B			Pf	PR1**DUP-02B			
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³		
Detections Reported by Laboratory	4	3	4	4	4	4		
Detections Impacted by Field Blank ⁴	4	3	3	4	2	4		
Usable Results ⁵	0	0	1	0	2	0		

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-17, Rev.3, July, 2008
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- 5. Based on Worksheet #11

The final recommended chlorinated herbicide sample collection method was impacted by field blank concentrations during Event #2, Attempt #2 for both the primary (HSM) and duplicate samples (LSM).

Table 13
Chlorinated Herbicides¹ – Target Analytes Impacted by Field Blank Concentrations by Collection
Method for Event #1, Attempt #3

	PR1CSOCLY**-01C			PF	R1**DUP-01C			
	ww	LSM ²	HSM ³	ww	LSM ²	HSM ³		
Detections Reported by Laboratory	4	4	4	4	4	4		
Detections Impacted by Field Blank ⁴	0	2	0	0	0	1		
Usable Results ⁵	4	2	4	4	4	3		

Notes:

- 1. Validation Guidance USEPA Region 2 SOP HW-17, Rev.3, July, 2008
- 2. LSM dissolved plus LSM particulate
- 3. HSM dissolved plus HSM particulate
- 4. Identified field blank contamination leading to positive results qualified as non-detect.
- Based on worksheet #11

The final recommended chlorinated herbicide sample collection method was impacted by field blank concentrations during Event #1, Attempt #3 for both the primary (HSM/whole water) and duplicate samples (LSM/whole water).

8. Cyanide

Cyanide is not a COPC/COPEC. The final recommended sample collection method selected was not impacted by field blank results because no positive results were "U" qualified on that basis. (Validation SOP reference: USEPA Region 2 SOP HW-2, Rev. 13, September, 2006.)

Appendix E - Field Blank Results Assessment

9. Volatile Organic Compounds

There are no COPC/COPECs in the TAL for VOCs. Field blank concentrations did not impact any result during Phase I for the VOCs identified in whole water or HSM sample collection methods. The final recommended sample collection method selected for VOCs was not impacted by field blank results.

10. Total Extractable Petroleum Hydrocarbons

TEPH is not a COPC/COPEC. Field blank results did not impact any TEPH result during Phase I in either the whole water or HSM sample collection methods. The final recommended sample collection method selected for TEPH was not impacted by field blank results. (Validation SOP reference: EDS SOP: TEPH-01, Rev.3, July, 2007).

Appendix F

Detailed Evaluation Sheets (Worksheet #11) – PCDDs/PCDFs

EVENT 2 ORIGINAL SAMPLE - DIOXIN PR1CSOCLY**-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

PCDD/PCDF Sample Collection Techniques	Sample Col	lection Qualit	ty ^a	Analytical Quality ^b	Identification of	Target Analyt	es
	Were specified sample meeting all analytical		ained	Are fewer than 2 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/ COPECs ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs ^c identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole water	No	Yes	NA	Yes	7	Yes	NA
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	4	Yes	NA
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	14	No	NA
LSM dissolved	No	Yes	NA	Yes	3	Yes	NA
HSM dissolved	No	Yes	NA	Yes	12	103	NA
LSM particulate	No	Yes	NA	Yes	4	Yes	NA
HSM particulate	No	Yes	NA	Yes	13	,,	NA

Positive Target Analyte Identification and Concentration Comparison^e

Analyte Identified	PR1CSOCLYWW-02B Whole Water ^f (pg/L)	LQ [€]	VQ	PR1CSOCLYLD-02B LSM Dissolved ^f (pg/L)	LQ ^g	VQ	PR1CSOCLYHD-02B HSM Dissolved ^f (pg/L)	LQ ^g	VQ	% RPD	PR1CSOCLYLP-02B LSM Particulate ^f (pg/g)	ΓŒ	VQ	PR1CSOCLYHP-02B HSM Particulate ^f (pg/g)	LQ ^g	VQ	% RPD
1,2,3,4,7,8-HxCDD	0.801	G		5000 017			0.606	c	3 J					6.32		J	
1,2,3,6,7,8-HxCDD	2.56	G					1.79	(S J		156	G		21.1		J	152
1,2,3,7,8,9-HxCDD	1.74	G	J	0.530	G	J	1.22	(5 J	78.9	114	G		15.2		J	153
1,2,3,4,6,7,8-HpCDD	84.3		J	11.0		J	38.5		J	111	4920			700		J	150
OCDD	1090			73.2		J	338		J	129	64000		j	9590	E	j	148
2,3,7,8-TCDF														3.82		М	
1,2,3,7,8-PeCDF														4.41	G	М	
2,3,4,7,8-PeCDF	0.537	G					0.288	(ã					4.04	G	М	
1,2,3,4,7,8-HxCDF							1.23	(i								
1,2,3,6,7,8-HxCDF							1.45	(ĵ.					11.7		М	
2,3,4,6,7,8-HxCDF	1.72	G					1.10	(ŝ					10.5		М	
1,2,3,4,6,7,8-HpCDF							17.3		J					205		J	
1,2,3,4,7,8,9-HpCDF							1.58	(3					13.3		J	
OCDF							42.3		ţ					444		j	

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

FFS = focused fesability study

HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCDD/PCDF = polychlorinated dibenzo-p-dioxin/polychlorinated dibenzofuran

pg/g = picoograms per gram

pg/L = picograms per liter

RPD = relative percent difference

VQ = validation qualifier - See Attachment 2 for definitions

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

COPCs/COPECs listed in the FFS: 2,3,7,8-TCDD; 1,2,3,7,8-PCDD; 1,2,3,6,7,8-HxCDD; 1,2,3,4,7,8-HxCDD; 1,2,3,7,8,9-HxCDD; 1,2,3,4,6,7,8-HpCDD; OCDD; 2,3,7,8-TCDF;

^{1,2,3,7,8-}PeCDF; 2,3,4,7,8-PeCDF; 1,2,3,6,7,8-HxCDF; 1,2,3,7,8,9-HxCDF; 1,2,3,4,7,8-HxCDF; 2,3,4,6,7,8-HxCDF; 1,2,3,4,6,7,8-HpCDF; 1,2,3,4,7,8,9-HpCDF; and OCDF.

^d Fewer than 2

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

f No rejected data.

⁸ A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 2 FIELD DUPLICATE - DIOXIN PR1**DUP-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

PCDD/PCDF	Camaria C	ollection Qua	i.aa	Analytical Quality ^b	Identification of	Tours Auglie	
nole water M dissolved plus HSM particulate M dissolved plus HSM particulate	Were specified san meeting all analytic	nple aliquots c		Analytical Quarity Are fewer than 2 results "R" qualified (rejected due to association with severe data quality issues)?		Are at least 2 more COPES' identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole water	No	Yes	NA	Yes	8	Yes	NA
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	11	Yes	NA
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	15	No	NA
LSM dissolved	No	Yes	NA	Yes	5	Yes	NA
HSM dissolved	No	Yes	NA	Yes	10		NA
LSM particulate	No	Yes	NA	Yes	9 11	Yes	NA
HSM particulate	No	Yes	NA	Yes	14	,,,,	NA

Positive Target Analyte Identification and Concentration Comparison^e

	PR1WWDUP-02B Whole Water ^f			PR1LDDUP-02B LSM Dissolved ^f			PR1HDDUP-02B HSM				PR1LPDUP-02B LSM Particulate ^f			PR1HPDUP-02B HSM Particulate ^f			
Analyte Identified	(pg/L)	LQ ^g	VQ	(pg/L)	LQ ^g	VQ	Dissolved ^f (pg/L)	LQ ^g	VQ	% RPD	(pg/g)	LQ ^g	VQ	(pg/g)	LQ ^g	VQ	% RPD
1,2,3,7,8-PeCDD											18.1	G		3.98	G	j	128
1,2,3,4,7,8-HxCDD	0.893	G	J	0.535	G	J	0.505	G	J	5.77				6.16		J	
1,2,3,6,7,8-HxCDD	2.76		J	0.548	G	J					106	G		19.8		J	137
1,2,3,7,8,9-HxCDD	1.94	G	J	100000			1.35	G	J		81.8	G		14.2		J	141
1,2,3,4,6,7,8-HpCDD	87.4		J	8.92		J	30.5		J	109	3160			636		J	133
OCDD	1230		J	64.7		J	199		J	102	43100			9560	E	J	127
2,3,7,8-TCDF														2.88		M	
1,2,3,7,8-PeCDF											11.1	G		4.04	G	M	93.3
2,3,4,7,8-PeCDF											11.8	O		4.23	G	М	94.4
1,2,3,4,7,8-HxCDF							0.959	G									
1,2,3,6,7,8-HxCDF	2.11	G					1.08	G			61.9	G		11.1		M	139
2,3,4,6,7,8-HxCDF	1.94	G					0.962	G			74.6	G		7.89		M	162
1,2,3,4,6,7,8-HpCDF							13.4		J					197		J	
1,2,3,4,7,8,9-HpCDF	2.61			0.515	G	J	1.20	G		79.9				12.5		J	
OCDF				1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			32.5		J					458		1	

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

FFS = focused fesability study

HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCDD/PCDF = polychlorinated dibenzo-p-dioxin/polychlorinated dibenzofuran

pg/g = picoograms per gram

pg/L = picograms per liter

RPD = relative percent difference

VQ = validation qualifier - See Attachment 2 for definitions

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

COPCs/COPECs listed in the FFS: 2,3,7,8-TCDD; 1,2,3,7,8-PeCDD; 1,2,3,6,7,8-HxCDD; 1,2,3,4,7,8-HxCDD; 1,2,3,7,8,9-HxCDD; 1,2,3,4,6,7,8-HpCDD; OCDD; 2,3,7,8-TCDF; 1,2,3,7,8-PeCDF; 2,3,4,7,8-PeCDF; 1,2,3,6,7,8-HxCDF; 1,2,3,7,8,9-HxCDF; 1,2,3,4,7,8-HxCDF; 2,3,4,6,7,8-HxCDF; 1,2,3,4,6,7,8-HyCDF; 1,2,3,4,7,8,9-HyCDF; and OCDF.

d Fewer than 2

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

f No rejected data.

⁸ A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

EVENT 1 ATTEMPT 3 ORIGINAL SAMPLE - DIOXIN PR1CSOCLY**-01C

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

PCDD/PCDF Sample Collection Techniques	Sample Co	ollection Qualit	v ^a	Analytical Quality ^b	Identification of	Target Analy	tes
	Were specified sampi all analytical needs?	le aliquots obta	ined meeting	Are fewer than 2 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/ COPECs ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole water	Yes	NA	Yes	Yes	14	No	No
LSM dissolved plus LSM particulate	Yes	NA	Yes	Yes	15	No	No
HSM dissolved plus HSM particulate	Yes	NA	Yes	Yes	15	No	No
LSM dissolved	Yes	NA	Yes	Yes	6	Yes	NA
HSM dissolved	Yes	NA	Yes	Yes	12	,03	NA
LSM particulate	Yes	NA	Yes	Yes	15	No No	No
HSM particulate	Yes	NA	Yes	Yes	15	,,,0	No

Positive Target Analyte Identification and Concentration Comparison^e

Analyte Identified	PR1CSOCLYWW-01C Whole Water ^f (pg/L)	LQ ^g	VQ	PR1CSOCLYLD-01C LSM Dissolved ^f (pg/L)	rde	vq	PR1CSOCLYHD-01C HSM Dissolved ^f (pg/L)	rđ	۷Q	% RPD	PR1CSOCLYLP-01C LSM Particulate ^f (pg/g)	LQ ^g	VQ	PR1CSOCLYHP-01C HSM Particulate ^f (pg/g)	rQ ^g	VQ	% RPD
1,2,3,7,8-PeCDD	0.425	G	J								24.4	G		4.56			137
1,2,3,4,7,8-HxCDD	0.914	G					0.575	G	J		47.7	G		9.01			136
1,2,3,6,7,8-HxCDD	2.58			0.769	G	1	1.42	G	J	59.5	135	G		24.4			139
1,2,3,7,8,9-HxCDD	2.01	G					1.04	G	J		105	G		17.5			143
1,2,3,4,6,7,8-HpCDD	81.5			13		J	31.3		J	82.6	3750		J	746			134
OCDD	1060		J	74.9		J	226		J	100	45500		1	12000	D		117
2,3,7,8-TCDF							0.0775	G			18.9	G		3.85			132
1,2,3,7,8-PeCDF	0.304	G				11	0.131	G	J		12.6	G		3,53			112
2,3,4,7,8-PeCDF	0.85	G									43.6	G		4.77			161
1,2,3,4,7,8-HxCDF	1.8	G					0.976	G	J		80.8	G		14.9			138
1,2,3,6,7,8-HxCDF	1.81	G		0.56	G	J	1.07	G	J	62.6	92.3	G		13.9			148
2,3,4,6,7,8-HxCDF	1.75	G		0.402	G	J	0.924	G	J	78.7	95.9	G		9,96			162
1,2,3,4,6,7,8-HpCDF	29.1		J	5.81		J	15.3		J	89.9	1760			253			150
1,2,3,4,7,8,9-HpCDF	2.05	G									105	G		13.8			154
OCDF	53.7		J				26.8		j		3280			488			148

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

FFS = focused fesability study

HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCDD/PCDF = polychlorinated dibenzo-p-dioxin/polychlorinated dibenzofuran

pg/g = picoograms per gram

pg/L = picograms per liter

RPD = relative percent difference

VQ = validation qualifier - See Attachment 2 for definitions

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPEcs listed in the FFS: 2,3,7,8-TCDD; 1,2,3,7,8-PCDD; 1,2,3,6,7,8-HxCDD; 1,2,3,4,7,8-HxCDD; 1,2,3,7,8,9-HxCDD; 1,2,3,4,6,7,8-HyCDD; 0CDD; 2,3,7,8-TCDF;

^{1,2,3,7,8-}PeCDF; 2,3,4,7,8-PeCDF; 1,2,3,6,7,8-HxCDF; 1,2,3,7,8,9-HxCDF; 1,2,3,4,7,8-HxCDF; 2,3,4,6,7,8-HxCDF; 1,2,3,4,6,7,8-HpCDF; 1,2,3,4,7,8,9-HpCDF; and OCDF.

^d Fewer than 2

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

[†] No rejected data.

g A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

EVENT 1 ATTEMPT 3 FIELD DUPLICATE - DIOXIN PR1**DUP-01C

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

PCDD/PCDF Sampl			_				
Collection Techniques Programmes	Sample Co	ollection Quali	tyª	Analytical Quality ^b	Identification	nof Target Ana	lytes
	Were specified sam meeting all analytica		ained	Are fewer than 2 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/ COPECs ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole water	Yes	NA	Yes	Yes	13	Yes	NA
LSM dissolved plus LSM particulate	Yes	NA	Yes	Yes	15	No	No
HSM dissolved plus HSM particulate	Yes	NA	Yes	Yes	15	No	No
LSM dissolved	Yes	NA	Yes	Yes	6	Yes	NA
HSM dissolved	Yes	NA	Yes	Yes	12	,,,,,	NA
LSM particulate	Yes	NA	Yes	Yes	15	No	No
HSM particulate	Yes	NA	Yes	Yes	15		No

Positive Target Analyte Identification and Concentration Comparison^e

	PR1WWDUP-01C			PR1LDDUP-01C LSM Dissolved ¹			PR1HDDUP-01C HSM Dissolved ^f				PR1LPDUP-01C LSM			PR1HPDUP-01C HSM Particulate ^f			
Analyte Identified	Whole Water ^f (pg/L)	LQ ^g	VQ	(pg/L)	LQg	VQ	(pg/L)	LQ ^g	VQ	% RPD	Particulate (pg/g)	LQ ^g	VQ	(pg/g)	LQ ^g	VQ	% RPD
1,2,3,7,8-PeCDD	0.262	G									59.3	G		4.69			171
1,2,3,4,7,8-HxCDD	0.681	G					0.448	G			91.2	G		9.24			163
1,2,3,6,7,8-HxCDD	1.81	G		0.652	G	J	1.18	G		57.6	219	G		25.0			159
1,2,3,7,8,9-HxCDD	1.30	G		0.419	G	J	0.834	G		66.2	238	G		21.0			168
1,2,3,4,6,7,8-HpCDD	71.1			10.4		j	29.3		J	95.2	7400		J	818			160
OCDD	821		J	72.8		J	269		J	115	109000		J	11600	D		162
2,3,7,8-TCDF							0.0948	G						3.60			
1,2,3,7,8-PeCDF											18.3	G		3.22			140
2,3,4,7,8-PeCDF	0.438	G									56.9	G		4.21			172
1,2,3,4,7,8-HxCDF	1.34	G		0.412	G		0.893	G		73.7	93.4	G		14.4			147
1,2,3,6,7,8-HxCDF	1.32	G					0.885	G			116	G		14.2			156
2,3,4,6,7,8-HxCDF	1.09	G					0.793	G			118	G		105			11.7
1,2,3,7,8,9-HxCDF				40							19.4	G					
1,2,3,4,6,7,8-HpCDF	20.2		J				13				2230			247			160
1,2,3,4,7,8,9-HpCDF	1.47	G		0.548	G	J	1.01	G		59.3	123	G		14.4			158
OCDF	38		J				23.1		J		4070			469			159

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern

 ${\sf COPECs = contaminants \ of \ potential \ ecological \ concern}$

FFS = focused fesability study

HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

 ${\tt PCDD/PCDF = polychlorinated\ dibenzo-p-dioxin/polychlorinated\ dibenzo furance}$

pg/g = picoograms per gram

pg/L = picograms per liter

 $\mathsf{RPD} = \mathsf{relative} \ \mathsf{percent} \ \mathsf{difference}$

 $VQ = validation\ qualifier - See\ Attachment\ 2\ for\ definitions$

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPECs listed in the FFS: 2,3,7,8-TCDD; 1,2,3,7,8-PeCDD; 1,2,3,6,7,8-HxCDD; 1,2,3,4,7,8-HxCDD; 1,2,3,7,8,9-HxCDD; 1,2,3,4,6,7,8-HpCDD; OCDD; 2,3,7,8-TCDF;

^{1,2,3,7,8-}PeCDF; 2,3,4,7,8-PeCDF; 1,2,3,6,7,8-HxCDF; 1,2,3,7,8,9-HxCDF; 1,2,3,4,7,8-HxCDF; 2,3,4,6,7,8-HxCDF; 1,2,3,4,6,7,8-HpCDF; 1,2,3,4,7,8,9-HpCDF; and OCDF.

d Fewer than 2

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

No rejected data.

⁸ A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Appendix G

Detailed Evaluation Sheets (Worksheet #11) – PCB Congeners

EVENT 2 ORIGINAL SAMPLE - PCB CONGENERS PR1CSOCLY**-02B

QAPP Worksheet #11-1

Project Quality Objectives/Systematic Planning Process Statements (Phase I)

Phase I Data Comparison Chart

PCB Congener Sample Collection Techniques	Sample Coll	lection Qualit	: y °	Analytical Quality ^b	ldent	tification of Target	t Analytes
	Were specified sample meeting all analytical r		ained	Are fewer than 17 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/COPECs ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCS/COPECS ^c identified (distinguished by asingle "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	No	Yes	NA	Yes	6	Yes	NA
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	7	Yes	NA
HSM dissolved plus HSM partiulate	No	Yes	NA	Yes	9	No	NA
LSM dissolved	No	Yes	NA	Yes	3	No	No
HSM dissolved	No	Yes	NA	Yes	2	140	No
LSIM particulate	No	Yes	NA	Yes	7	Yes	NA
HSM particulate	No	Yes	NA	Yes	9	162	NA

	PR1CSOCLYWW-02B			PR1CSOCLYLD-02B LSM Dissolved ^E			PR1CSOCLYHD-02B HSM Dissolved ⁶				PR1CSOCLYLP-02B LSM Particulate ⁸			PR1CSOCLYHP-02B HSM Particulate ^g		
Analyte Identified	Whole Water ^g (pg/L)	ŁQ ^h	VQ	(pg/L)	ľď,	VQ	(pg/L)	LQh	VQ	% RPD	(pg/g)	LQ"	VQ	(pg/g)	LQ" VQ	% RPD
PCB-1	14.1	D		13.4	D		18.4	D		31.4				204	D M	
PCB-4/10														915	DM	
PCB-6	26.6	D		13.6	DG		25.3	D		60.2				446	DM	
PCB-16/32														1840	D M	
PCB-17						1000								1250	DМ	
PCB-18														2590	D M	
PCB-19	28.3	D												420	DM	
PCB-22														1140	DJ	
PCB-25														480	L Q	
PCB-26														701	DJ	
PCB-28				5.0										3310	DJ	
PCB-31														2970	DJ	
PCB-35				3.63	DG		7.07	DG		64.3	879	DG		204	DM	125
PCB-36											478	DG		98.6	DG M	132
PCB-40														718	DM	
PCB-41/64/71/72														3360	DM	
PCB-42/59							100000							1210	DJ	
PCB-43/49														2970	D J	
PCB-44														3890	DM	
PCB-45														611	DM	
PCB-46	9.49	DG		3.20	DG		9.59	DG		99.9	848	DG		303	DM	94.7
PCB-48/75	22.3	D												677	DM	
PCB-52/69														4780	DJ	
PCB-53														596	DM	
PCB-55														90.2	DG M	
PCB-56/60														2400	DM	
PCB-57														26.9	DG M	
PCB-58														16.6	DG M	
PCB-61/70														4540	DJ	
PCB-63					0.00		4.20	DG			497	DG		153	D J	106
PCB-67											383	DG		113	DM	109
PCB-74														1450	L Q	
PCB-76/66														3020	LIQ.	
PCB-79				1.92	DG						420	DG				
PCB-81											450	DG				
PCB-82	46.1	D	J											1170	ΙŪ	
PCB-84/92	129	D	J											3580	D M	
PCB-85/116	48.9		J	10.5	D		25.6	D		83.7				1400	DM	
PCB-87/117/125	117	D												3400	DM	
PCB-88/91	40.6		j	200						l				1060	DJ	
PCB-89	1	_	1			1000					393	DG		90.3	DG M	125
PCB-90/101	309	D	J				189	D						8320	DM	
PCB-94	1		1											37	DG J	
PCB-95/98/102	211	D	J											5790	DJ	
PCB-96	1		İ											65.5	DG J	
PCB-97	95.4	D	ı j											2490	DM	
PCB-99	114		ı j				66.0	D						3280	DM	
PCB-100	124		Ė											27.6	DG J	
PCB-103	1		†											52.5	DJ	
PCB-105	122	D	di .											3350	D M	
PCB-106/118	269	D								-				7890	DM	
PCB-107/109	20.4	D		4.71	DG	1027	10.8	D		78.5				503	DJ	
PCB-107/109 PCB-108/112	15.8	D		3.54	DG		9.84	DG		94.2	1110	DG		403	DM	93.5
PCB-110	353	D		3,34	DG		9.84	DG		34.2	1110	UG		9800	DM	95.5
I CD-110	353	<u>u</u>	13				3.66	DG			544	DG		183	DM	99.3

	PR1CSOCŁYWW-02B			PR1CSOCLYLD-02B LSM Dissolved ⁸			PRICSOCLYHD-02B				PRICSOCLYLP-02B			PR1CSOCLYHP-02B HSM Particulate ⁸			
Analyte Identified	Whole Water ^g (pg/L)	ŁQ ^b	VQ	(pg/L)	ro,	VQ	(pg/L)	LQ ^h	VQ	% RPD	(pg/g)	LQb	VQ	(pg/g)	ια	VQ	% RPD
PCB-114	6.37	DG	J	1.96	DG						430	DG		175		D M	84.3
PCB-119	7.55	DG	J				2.64	DG			424	DG		142		D M	99.6
PCB-122				1.21	DG						261	DG		88.5		G M	98.7
PCB-123											590	DG		148		D J	120
PCB-124	14.5	D	j	3.24	DG		7,76	DG		82.2	978	DG		379		D J	88.3
PCB-126											529	DG		82.1		gМ	146
PCB-128/162	60.0	D	J				27.8	D	j		4830	DG		1880	A DESCRIPTION OF THE PERSON OF	D M	87.9
PCB-129	20.0	D	J	4.36	DG		9.54	DG	j	74.5	1330	DG		590		DМ	77.1
PCB-130	20.0	D	J	5.11	DG		10.4	D	j	68.2	1890	DG		666		οм	95.8
PCB-132/161	90.7	D	j				42.2	D						2890		D M	
PCB-133/142	11.1	D	J	2.27	DG						778	DG		304		D M	87.6
PCB-134/143	18.4	D	j	4,27	DG									537		D M	
PCB-135	40.1	D		8.76	DG		19.9	D		77.7				1180		DМ	
PCB-136														1110		D M	
PCB-137	17.7	D	J	3.88	DG		11.7	D	1	100				460		οм	
PCB-138/163/164	334	D		5,00	93		162	D		<u> </u>				10100		D M	
PCB-139/149	210	D					102							6730		D M	
PCB-141	59.9	D		0.55										1870		DM	
PCB-144	33.3		•	3.39	DG		8.35	DG		84.5	1380	DG		448		D M	102
PCB-146/165	38.3	D	1	-			1							1140		DМ	
PCB-147	36.3	۲	,								679	DG		170		D M	120
PCB-151											0,3	-		1850		D M	120
PCB-153	265	D				1000000								7950		D M	
PCB-154	203		,											74.8		G M	
PCB-155							3.19	DG						74.0	-	G IVI	
PCB-156	37.4	D	1				3,13	00						1070	١.,	D M	
PCB-157	11.7	D		2.30	DG		4,94	DG	1	72.9	1020	DG		269		D M	117
PCB-158/160	39.1	D		2.50	Du		4,54			12.5	1020	- 50		1220		D M	
PCB-166	33.1	۲	,											59.5		D M	
PCB-167	14.5	D	1	3.51	DG		7.68	DG		74.5	1110	DG		436		D M	87.2
PCB-168	14.5	۲	,	3.31	20		7.00	- 00	•	14.5	1110	- 00		7.35		G M	- 07.Z
PCB-170	72.1	D												2600		D M	
PCB-171	22.3	D												658		D M	
PCB-172	15.3	D		3.55	DG		7.64	DG	1	73.1	1210	DG		444		D M	92.6
PCB-173	13.3		J	رد.د	<i>D</i> 6		7.04	DG	•	/3.1	1210	UG		69.9	O CHARLEST AND ADDRESS.	G M	92.0
PCB-174										-				2470		D M	
PCB-175														116		D M	
PCB-176														320		D M	
PCB-177	43.3	D								-				1500		D M	
PCB-178	17.8	D		4.67	DG						1530	DG		552		D M	93.9
PCB-179	17.0	-	,	4.07	ы						1330	DG		1150		D M	93.5
PCB-180														5600		D M	
PCB-180 PCB-182/187														3410	ALTONOOPING STATE	D M	
PCB-183														1440		D M	
PCB-183				2.40	DG		7.92	DG	1	107	470	DG		1440		IVI	
PCB-184 PCB-185	1			2.40	DG		7.92	UG	y	10/	470	DG		317	—	D M	_
				2.01	DG					-	483	DG		116		D M	122
PCB-189	_									-	483	DG		116 468		D M	123
PCB-190	1			1.36	DG					-				93.1		G M	\vdash
PCB-191	2.10	5.0		1.25	DG DG		3.00	5.0		61.0							00.1
PCB-193	9.42	DG	J	2.10	DG		3.98	DG	1	61.8	711	DG		283		D J	86.1
PCB-194						90 000								1580		CONTRACTOR OF THE PARTY OF	
PCB-195	15.8	D	J				6.95	DG	ı	-	1180	DG		647		DJ	58.3
PCB-196/203														1840		D M	
PCB-198										-				78.3		G M	1
PCB-199	42.0	D	J											1940		οм	

Analyte Identified	PR1CSOCLYWW-02B Whole Water ^g (pg/L)	LQ ^h	VQ	PR1CSOCLYLD-02B LSM Dissolved ⁱⁱ (pg/L)	LQ ^h	VQ	PR1CSOCLYHD-02B HSM Dissolved ^E (pg/L)	ια ^h	VQ	% RPD	PR1CSOCLYLP-02B LSM Particulate ^s (pg/g)	ιQ ^h	VQ	PR1CSOCLYHP-02B HSM Particulate [®] (pg/g)	rā, kā	% RPD
PCB-200						10.00								203	D M	
PCB-201											517	DG		230	DM	76.8
PCB-202											934	DG		450	DM	69.9
PCB-206														2250	DJ	
PCB-207														238	DJ	
PCB-208							100							749	DJ	

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern FFS = focused fesability study HSM = high-solids mass LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCB = polychlorinated biphenyl pg/g = picograms per gram

pg/L = picograms per liter RPD = relative percent difference

VQ = validation qualifier - See Attachment 2 for definitions

^b Analytical quality is based upon the program 90% analytical completeness objectives.

[°] COPCs/COPECs listed in the FFS: PCB-77, PCB-81, PCB-105, PCB-114, PCB-118, PCB-123, PCB-126, PCB-156, PCB-157, PCB-167, PCB-169, and PCB-189.

d At least 2

e Fewer than 17

Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

⁸ No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 2 FIELD DUPLICATE - PCB CONGENERS PR1**DUP-02B

QAPP Worksheet #11-1

Project Quality Objectives/Systematic Planning Process Statements (Phase I)

Phase I Data Comparison Chart

PCB Congener Sample Collection Techniques	Sample Co	illection Qual	ity ^a	Analytical Quality ^b	ldent	ification of Targe	t Analytes
	Were specified sam meeting all analytic		btained	Are fewer than 17 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCS/COPECs ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs ^c identified (distinguished by asingle "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	No	Yes	NA	Yes	7	Yes	NA
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	8	No	No (62)
HSM dissolved plus HSM partiulate	No	Yes	NA	Yes	9	No	Yes (138)
LSM dissolved	No	Yes	NA	Yes	3	No	No
HSM dissolved	No	Yes	NA	Yes	3	,40	No
LSM particulate	No	Yes	NA	Yes	8	No	Yes
HSM particulate	No	Yes	NA	Yes	9	NO	Yes

	PR1WWDUP-02B			PR1LDDUP-02			PR1HDDUP-02B				PR1LPDUP-02B			PR1HPDUP-02B			
	Whole Water ^g	h		LSM Dissolve			HSM Dissolved ⁸				LSM Particulate ⁸	6		HSM Particulate ⁸	- 11	1	
Analyte Identified	(pg/L)	LQ ^h	VQ	(pg/L)	LQh	VQ	(pg/L)	LQ	VQ	% RPD	(pg/g)	rd,	VQ	(pg/g)	rđ	NAME OF TAXABLE PARTY.	% RPD
PCB-1	19.9	D		1	i.7 [)	19.3	D		14.4				192		M	
PCB-4/10														1080	2000	M	
PCB-6	27.0	D		1	.1 DG		25.7	D		52.0				639		M	
PCB-15														1430		М	
PCB-16/32														2250		M	1
PCB-17														1670		М	ĺ
PCB-18														2970		M	
PCB-19	25.3	D												564		M	
PCB-20/21/33					2									2230		M	
PCB-22														1960	D		
PCB-25	24.8	D												4100	۵		
PCB-26														2680	D	regardous gar	
PCB-28														15100	D		
PCB-31									(0)					9100	D	The strategy	
PCB-35	8.56	DG					5.95	DG						242	D	M	
PCB-36											291	DG					Į.
PCB-37														2050	D		
PCB-40														1030		M	0
PCB-41/64/71/72														5090		M	1000
PCB-42/59						100								2380	D		
PCB-43/49						100								9130	D	J	
PCB-44														6390		M	
PCB-45														755	۵	M	
PCB-46	12.3	D		4	28 DG		10.2	D	1	81.8	610	DG		450		M	30.2
PCB-47														5580	D	J	-
PCB-48/75	24.4	D												1110	D	M	
PCB-51														522	D	J	
PCB-52/69														8660	D	J	
PCB-53									100					966	D	M	
PCB-54														47.2	D	M	
PCB-55	3.56	DG												103	D	M	4.00
PCB-56/60														3320	D	M	
PCB-57														49.0	D	M	
PCB-61/70	172	D												7700	D	j.	
PCB-63	5.64	DG									346	DG		670	D	j	63.8
PCB-67	3.19	DG												240	A 100	M	
PCB-68																	
PCB-74														3490	D	j i	
PCB-76/66	118	D	J											7430	D	J	
PCB-77																	
PCB-79	3.49	DG		1	42 DG												
PCB-81											88.9	DG					Į.
PCB-82	42.0	D									2770			1470	D	M	61.3
PCB-84/92	114					1000					n in the second			4720		M	

	PR1WWDUP-02B			PR1LDDUP-02B			PR1HDDUP-02B				PR1LPDUP-02B			PR1HPDUP-028		
Analyte Identified	Whole Water ^g (pg/L)	LQ ^ħ	VQ	LSM Dissolved ⁸ (pg/L)	LQh	VQ	HSM Dissolved g (pg/L)	LQt	VQ	% RPD	LSM Particulate ⁸ (pg/g)	LQ"	vq	HSM Particulate ⁸ (pg/g)	LQ" VQ	% RPD
PCB-85/116	47.1	D		11.4	D		24,1	D	1	71.5				1760	DM	
PCB-87/117/125	113	D									7180	D		4290	D M	50.4
PCB-88/91	37.0	D												1510	DM	
PCB-89											275	DG		129	D M	72.3
PCB-90/101	283	D			100		193	D						11200	DM	
PCB-95/98/102	200	D												7820	DM	
PCB-96	2.08	DG														
PCB-97	86.8	D												3250	D M	
PCB-99	112	D					66.3	D	J					4780	D M	
PCB-103	1.74	DG		1000		10000										
PCB-105	104	D									7470	D		4050	DM	59.4
PCB-106/118	266	D					144	D	J		16800	D		10500	DM	46.2
PCB-107/109	16.2	D		4.94	DG		10.9	а	J	75.3	1330	DG		750	DM	55.8
PCB-108/112	13.4	D		3.30	DG		8.68	DG	j	89.8	980	DG		524	DM	60.6
PCB-110	307	D												12300	DM	
PCB-111/115	3.75	DG		1.77	DG						398	DG		192	DM	69.8
PCB-114	6.56	DG		1.18	DG						471	DG		213	D M	75.4
PCB-119	5.01	DG					3.14	DG	ı		353	DG		240	D M	38.1
PCB-122														110	D M	
PCB-123							4,52	DG	j		432	DG		179	DM	82.8
PCB-124	12.7	D					7.85	DG			850	DG		464	D M	58.8
PCB-126	3.72	DG												95.7	DM	
PCB-128/162	55.3	D					27.6	D	j		4140	D		2320	D M	56.3
PCB-129	19.7	D		4,58	DG		10.8	D	j.	80.9	1290	DG		741	D M	54.1
PCB-130	19.9	D		4.45	DG		10.9	D	J	84.0	1560	DG		868	D M	57.0
PCB-132/161	85.6	D					47.0	D	ı		6000	D		3480	DM	53.2
PCB-133/142	8.92	DG		2.27	DG		5.73	DG	1	86.5	597	DG		374	D M	45.9
PCB-134/143	17.6	D		4.21	DG									689	DM	
PCB-135	41.0	D		8.82	DG		20.3	D	J	78.8				1520	DM	
PCB-136	34.5	D												1460	DM	
PCB-137	13.7	D		3.78	DG		12.9	D	ı	109				665	D M	
PCB-138/163/164	313	D					166	D	J		20800	D		12300	D M	51.4
PCB-139/149	206	D												8730	D M	
PCB-140											215	DG				
PCB-141	62.9	D												2340	DM	
PCB-144	11.9	D		3.98	DG		7.19	DG	J	57.5	852	DG		507	D M	50.8
PCB-146/165	34.6	D												1400	D M	
PCB-147														270	DM	
PCB-151														2250	D M	
PCB-153	243	D												9230	DM	
PCB-154														123	D M	
PCB-155	2.78	DG		1.40	DG		3.26	DG	J	79.8						
PCB-156	30.5	D									2280	DG		1350	D M	51.2
PCB-157	7.79	DG		2.49	DG						720	DG		354	D M	68.2
PCB-158/160	36.4	D												1520	D M	
PCB-166														51.9	D M	
PCB-167	13.8	D		2.85	DG		6.65	DG	J	80.0	968	DG		537	D M	57.3
PCB-170	72.1	D									5490	D		2800	D M	64.9
PCB-171	20.4	D									1560	DG		716	D M	74.2
PCB-172	12.9	D		3.44	DG		7.93	DG	1	79.0	1060	DG		505	D M	70.9
PCB-174														2680	D M	

Analyte Identified	PR1WWDUP-02B Whole Water ^g (pg/L)	LQ ^h	VQ	PR1LDDUP-02B LSM Dissolved ⁸ (pg/L)	LQ ^h	VQ	PR1HDDUP-02B HSM Dissolved ^g (pg/L)	ια'	VQ	% RPD	PR1LPDUP-02B LSM Particulate ⁸ (pg/g)	LQ ^h	PR1HPDUP-02B HSM Particulate ⁸ VQ (pg/g)	IQ ^h VQ	% RPD
PCB-175													137	DM	
PCB-176													352	DM	
PCB-177	41.1	D									2990	DG	1590	DM	61.1
PCB-178	17.9	D		3.77	DG		9.17	DG	j	83.5	1180	DG	653	DM	57.5
PCB-179													1250	DM	
PCB-180				4.000000000									6220	DM	
PCB-182/187													3790	DM	
PCB-183													1710	DM	
PCB-184	7.15	DG		1.87	DG						291	DG			
PCB-185	9.38	DG					5.10	DG	J		725	DG	333	DM	74.1
PCB-189				2.000									118	DM	
PCB-190													552	DM	
PCB-191	3.07	DG											113	DM	
PCB-193	6.51	DG		1.81	DG						503	DG	276	DM	58.3
PCB-194				30.55.34									1480	DM	
PCB-195	13.8	D	J				8.01	DG	J				707	D M	
PCB-196/203													1820	DM	
PCB-197													66.9	DG M	
PCB-198													95.2	DG M	
PCB-199	36.6	D		8.24	DG								1750	DM	
PCB-200													242	D M	
PCB-201	5.85	DG									506	DG	227	DM	76.1
PCB-202	11.1	D		2.41	DG						765	DG	410	D M	60.4
PCB-206													1420	DJ	
PCB-208													441	l d	

a Na" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study
HSM = high-solids mass

LSM = low-solids mass $LQ = laboratory qualifier - See Attachment 1 for definitions <math display="block">PCB = polychlorinated biphenyl \\ pg/g = picograms per gram$

pg/L = picograms per liter RPD = relative percent difference VQ = validation qualifier - See Attachment 2 for definitions % = percent

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

[°] COPCs/COPECs listed in the FFS: PCB -77, PCB -81, PCB -105, PCB -114, PCB -118, PCB -123, PCB -126, PCB -156, PCB -157, PCB -167, PCB -169, and PCB -189.

d At least 2

e Fewer than 17

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

⁸ No rejected data.

^h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 1 ATTEMPT 3 ORIGINAL SAMPLE - PCB CONGENERS PR1CSOCLY**-01C

QAPP Worksheet #11-1

Project Quality Objectives/Systematic Planning Process Statements (Phase I)

Phase I Data Comparison Chart

PCB Congener Sample Collection Techniques	Sample Colle	ction Quality	а	Analytical Quality ^b	ldent	tification of Target	Analytes
	Were specified sample a all analytical needs?	fliquots obtain	ned meeting	Are fewer than 17 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/COPECs ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs ^c identified (distinguished by asingle "no" in the previous column), are the overall number of target a significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	Yes	NA	Yes	Yes	6	Yes	NA
LSM dissolved plus LSM particulate	Yes	NA	Yes	Yes	8	No	No (120)
HSM dissolved plus HSM partiulate	Yes	NA	Yes	Yes	9	No	Yes (153)
LSM dissolved	Yes	NA	Yes	Yes	2	Yes	NA
HSM dissolved	Yes	NΑ	Yes	Yes	6	, 63	NA
LSM particulate	Yes	NA	Yes	Yes	8	No	No
HSM particulate	Yes	NA	Yes	Yes	8	NO	Yes

Analyte Identified	PR1CSOCLYWW-01C Whole Water ^g (pg/L)	ì.Q ^ħ	VQ	PR1CSOCLYLD-01C LSM Dissolved ⁸ (pg/L)	rđ _r	VQ	PR1CSOCLYHD-01C HSM Dissolved [®] (pg/L)	LQ"	VQ	% RPD	PRICSOCLYLP-01C LSM Particulate ^E (pg/g)	10 ^h	va	PR1CSOCLYHP-01C HSM Particulate ⁶ (pg/g)	TO,	VQ	% RPD
PCB-1	vinole viate: (pg/c/	1	1	======================================		1	11-07-1	1	1	70.111.0	(Para)		1	177	STREET	o on second	
PCB-4/10	135	D	1	120	n	-		1						1550	D, G		
PCB-5/8	133	<u> </u>		110										2190	D		
PCB-6						1		-						810	D	A	
PCB-11								†					-	5120	GEODESIUS)	1	
PCB-15														779	D	700000000000000000000000000000000000000	
PCB-16/32														2920		J	
PCB-17	130	D	J											2450	D	The second second	
PCB-18														2820	D		
PCB-19	53.8	D	J											827	D	li i	
PCB-20/21/33														1670	D	J	
PCB-22														1710	D	j	
PCB-24/27														467	D	-	
PCB-25	41.4	D	j											919	D	J	
PCB-26														1080	D	J	
PCB-28														5920	D	j	
PCB-31														4580	D	j	
PCB-35	11.2	D					4.08	D,G			1540	D	1	267	D	1	141
PCB-37														1620	D	J	
PCB-40											7030	D	J	1080	D	J	147
PCB-41/64/71/72	149	B, D	J								31700	B, D	J	5330	D) J	142
PCB-42/59	62.3	D	J								11100	D	J	1990	D	J	139
PCB-43/49	163	D	j								34100	B, D	1	5450	D	J	145
PCB-44	179	B, D	J								34400	B, D	J	5720	٥	J	143
PCB-45											5830	D	J	767	D	J	153
PCB-46	20.1	D									3550			523	D	. J	149
PCB-47											14400		J	2690	D	J	137
PCB-48/75							100				6340		J	685	D	J	161
PCB-50	14.1	D		4.91	D, G		8.70	D, G		55.7	1300						
PCB-51											2900			560	D		135
PCB-52/69		B, D	J								45200		1	6570	D		149
PCB-53	43.9	D	J								6630	D	J	1170	D		140
PCB-55														130	D, G		
PCB-56/60											27600		J	4400	D	A LONG LAND CO.	145
PCB-61/70	200		J								45500		J	6590	D	The second second	149
PCB-63		D, G									1950	D		330	D		142
PCB-67		D, G												153	D, G		
PCB-74	61.0		J								16800	COMPLETE STATE	J	2340	D		151
PCB-76/66	150	D	J		10 G 10						35700		I	6080	D	0.001.001.000.00	142
PCB-77											4370	D	J	856	D	A COLUMN TO SERVICE	134
PCB-79	3.01	D, G												146	D, G	1	
PCB-81						10.00		D, G									
PCB-82	45.6	D	J	11.5	D		18.4	D	J	46.2	8130	D	1	1550	D	J	136
PCB-83		1	1														A

							PR1CSOCLYHD-01C		1		PRICSOCLYLP-01C			PR1CSOCLYHP-01C			
	PR1CSOCLYWW-01C			PR1CSOCLYLD-01C			HSM Dissolved ⁸				LSM Particulate ^g			HSM Particulate ⁸			á
Analyte Identified	Whole Water ^g (pg/L)	LQ ^b	VQ	LSM Dissolved [®] (pg/L)	LQ ^h	VQ	(pg/L)	LQ*	VQ	% RPD	(pg/g)	LQ ^h	VQ	(pg/g)	LQ ^b	VQ	% RPD
PCB-84/92	129	B, D	j				1000000				23700	D	J	4010	D	J	142
PCB-85/116	47.1	D	J	14.1	D		21.9	D	J	43.3	9720	D	J	1980	D	j	132
PCB-87/117/125	121	D	j				50.6	D	J		19800	D	J	3780	D	J	136
PCB-88/91	40.3	D	j	13.0	D		21.9	D	j .	51.0	8370	D	J	1380	D	J	143
PCB-89							1.14	D, G	ı								
PCB-90/101	288	B, D	j	77,8	D						49600	B, D	J	8740	D	j	140
PCB-95/98/102	221	B, D	J								37800	B, D	J	6140	D	j	144
PCB-96																	
PCB-97	90.7	D	J								15900	D	J	3050	D	J	136
PCB-99	116	B, D	J				52.6	D	J		21700	D	J	4060	D	J	137
PCB-105	113	D	J			70.00	44.6	D			18300	D	J	4080	D	J	127
PCB-106/118	266	B, D	J				123	B, D	j.		46900	B, D	j	9370	D	J	133
PCB-107/109	19.6	D					8.47	D, G	J		3600	D		748	D	J	131
PCB-108/112	15.1	D					7.53	D, G	J		2910	D		494	D	J	142
PCB-110	343	B, D	J				149	B, D	ı		59600	B, D	J	11400	D	j	136
PCB-111/115	5.52	D, G									1490	D		202	D, G	J	152
PCB-114	5.85	D. G									1400	D		208	D, G	j	148
PCB-119	5.70				700	100								178	D, G		Å
PCB-124	13.2	D					5.52	D. G	j.		2360	D		475	D	J	133
PCB-126					1000000									130	D, G	j	
PCB-128/162	62.5	D	j	13.6	D		21,9	D		46.8	9740	D	J	2110	D	j	129
PCB-129	23.0	D		4.18	D, G		7.82	D, G		60.7	3070	D		636	D	j	131
PCB-130	22.5	D	J	5.46	D. G		7.45	D, G		30.8	3500	D		757	D	j	129
PCB-132/161	97.6	D	j								14000	D	J	3090	D	J	128
PCB-133/142	10.1	D									1790	D		309	D	J	141
PCB-134/143	18.0	D		4.56	D, G		7.02	D, G		42.5	2820	D		611	D	J	129
PCB-135	50.1	D	J	12.7	D		19.1	D	J.	40.3	9070	D	J	1350	D	J	148
PCB-136	41.7	D	J	12.2	D		21.6	D	J.	55.6	7700	B, D	J	1180	D	1	147
PCB-137	18.0	D		4.37	D, G		8.13	D, G		60.2	3500	D		634	D	J	139
PCB-138/163/164	365	B, D	J				126	B, D			56500	B, D	J	11700	D	J	131
PCB-139/149	267	D	J	76.4	D		114	D	J.	39.5	51100	B, D	ı	8060	D	J	146
PCB-141	71.8	D	J								12400	D	J	2240	D	J	139
PCB-144	16.1	D									3280	D		477	D	J	149
PCB-146/165	40.9	D	J								6530	D	J	1240	D	J	136
PCB-147	7.99	D, G												216	D	J	
PCB-151	71.6	B, D	j	19.6	D						15500	B, D	J	2100	D	J	152
PCB-153	286	B, D	j	100			108	B, D			50400	B, D	J	9110	D	J	139
PCB-155	4.23	D, G															
PCB-156	39.1	D	j	6.64	D,G		12.8	D		63.4	6020	D	J	1250	D	J	131
PCB-157	9.10	D, G					3.71	D, G			1550	D		336	D	J	129
PCB-158/160	44.7	D	J								6810	D	J	1410	D	j	131
PCB-167	15.8	D		3.65	D, G		5.34	D, G		37.6	2430			527	D	The second	129
PCB-170	99.9	D	J	20.6	D		31.1	D		40.6	17600	D	J	2900	D	J	143
PCB-171	26.0	D	J	5.71			8.38	D, G		37.9	4560	D	J	826	D	J	139
PCB-172	17.1	D	j	3.69	D, G		6.64	D, G		57.1	3370	D		589	D	J	140

Analyte Identified	PR1CSOCLYWW-01C Whole Water ^s (pg/L)	ŁQ ^h	VQ	PR1CSOCLYID-01C LSM Dissolved [®] (pg/L) LQ ^N	VQ	PR1CSOCLYHD-01C HSM Dissolved ^a (pg/L)	LQ ^b	VQ	% RPD	PRICSOCLYLP-01C LSM Particulate ⁸ (pg/g)	LQ'n	VQ	PR1CSOCLYHP-01C HSM Particulate [®] (pg/g)	ŁQ ^h	VQ	% RPD
PCB-174	104	D j		21.9 D		31.6	D		36.3	18500	D	j	3010	D	J	144
PCB-175										1070	D, G		104	D, G	J	165
PCB-176	13.1	D		3.19 D, G		5.02	D, G		44.6	2560	D		354	D	J	151
PCB-177	60.8	D j		11.2 D		19.1	D		52.1	10200	D	1	1700	D	J	143
PCB-178				5.43 D, G		9.00	D, G		49.5	5090	D	J	719	D	J	150
PCB-179	47.0	D J		0.000						9850	D	1	1320	D	J	153
PCB-180	222	B, D J								42700	D	J	6910	D	J	144
PCB-182/187	133	D J		29.8 D		47.8	D		46.4	30800	D	J	4150	D	ı J	153
PCB-183	60.7	D J		13.4 D		20.1	D		40.0	12400	D	j	1890	D	J	147
PCB-184				3.24 D, G		6.67	D, G		69.2	805	D, G					50000
PCB-185	13.0	D		3.28 D, G		5.04	D, G		42.3	2500	D		361	D	J	150
PCB-189				Harden and the second	100000					717	D, G					
PCB-190	19.1	D J		4.22 D, G		6.10	D, G		36.4	3410	D		585	0	J	141
PCB-191										851	D, G		129	D, G	J	147
PCB-193	8.85	D, G		2.26 D, G		3.49	D, G		42.8	1960	D		309	D	ı J	146
PCB-194	49.2	D j		8.82 D, G		14.7	D		50.0	11200	D	1	1710	D		147
PCB-195	21.8	D j		3.90 D, G						4570	D	J	667	D		149
PCB-196/203	54.5	D j		13.0 D		23.0	D		55.6	18400	D	1	1900	D	J	163
PCB-199	53.0	D j		12.2 D		19.4	D		45.6	18800	D	j	1870	D	J	164
PCB-200	7.49	D, G								2680	D		263	D) J	164
PCB-201	8.62	D, G				3.30	D, G			2300	D		244	D	J	162
PCB-202	15.0	D j		3.78 D, G		5.38	D, G		34.9	3900	D	J	414	D	J	162
PCB-206	35.6	D J		200						8100	D	j	1430	D		140
PCB-207	3.87	D, G J								941	D, G					o de la companya de l
PCB-208	11.5	D J			10000	3.26	D, G			2590	D		498	D		135
PCB-209													1130	D		

a Na" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern FFS = focused fesability study HSM = high-solids mass LSM = low-solids mass $LQ = laboratory qualifier - See Attachment 1 for definitions \\ PCB = polychlorinated biphenyl \\ pg/g = picograms per gram$

pg/L = picograms per liter RPD = relative percent difference VQ = validation qualifier - See Attachment 2 for definitions % = percent

^b Analytical quality is based upon the program 90% analytical completeness objectives.

[°] COPCs/COPECs listed in the FFS: PCB -77, PCB -81, PCB -105, PCB -114, PCB -118, PCB -123, PCB -126, PCB -156, PCB -157, PCB -167, PCB -169, and PCB -189.

d At least 2

e Fewer than 17

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

g No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 1 ATTEMPT 3 FIELD DUPLICATE - PCB CONGENERS PR1**DUP-01C

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

PCB Congener Sample Collection Techniques	Sample Col	lection Quali	ty ^a	Analytical Quality ^b	lden	tification of Targe	t Analytes
	Were specified sampl meeting all analytical		tained	association with severe data	Number of COPCs/COPECs ^C listed in the FFS identified?	Are at least 2 more COPCs/COPECs ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by asingle "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	Yes	NA	Yes	Yes	6	Yes	NA
LSM dissolved plus LSM particulate	Yes	NA	Yes	Yes	5	Yes	NA
HSM dissolved plus HSM partiulate	Yes	NA	Yes	Yes	9	No	NA
LSM dissolved	Yes	NA	Yes	Yes	3	Yes	NA
HSM dissolved	Yes	NA	Yes	Yes	5	les les	NA
LSM particulate	Yes	NA	Yes	Yes	5	Yes	NA
HSM particulate	Yes	NA	Yes	Yes	9	165	NA

				PR1LDDUP-01C			PR1HDDUP-01C				PR1LPDUP-01C			PR1HPDUP-01C			
Analyte Identified	PR1WWDUP-01C Whole Water ^s (pg/L)	LQ ^h	VQ.	LSM Dissolved ^g (pg/L)	LQ ^b	VQ	HSM Dissolved ⁶ (pg/L)	LQ ^h	vq	% RPD	LSM Particulate ⁸ (pg/g)	LQ ^h	VQ	HSM Particulate ^g (pg/g)	LQ ^h	νq	% RPD
PCB-1	1			,,,,				•			,, 0, 0,			161	SHOW BOOK INC.	0.000	
PCB-4/10	170	D	J	129	D		-500				2600	D	1	1420	D		58.7
PCB-5/8				-							-			1970	D	Selection of the selection of	
PCB-6														806	D		
PCB-7/9							7.65	D, G									
PCB-11														4130	D		
PCB-15														819	D		
PCB-16/32	259	D	J		30.00									3680	D		
PCB-17	226	D	J											3360	Q		
PCB-18														3560	D		
PCB-19	85.9	D	J											933	D		
PCB-20/21/33								100						1170	D		
PCB-22														1100	D		
PCB-24/27	41.6	D												605	D		
PCB-25	66.6	D	J											1060	D		
PCB-26	70.9	D	J											950	D		
PCB-28	344	D	J											4500	D		
PCB-31														3710	D		
PCB-35	17.0	D					3.96	D, G	j					211	D		
PCB-37														1070	D		
PCB-40	48.1	D												771	D		
PCB-41/64/71/72	238	B, D	J											3960	D		
PCB-42/59	95.9	D	J											1470	a		
PCB-43/49	279	D	J											4130	D		
PCB-44	279	B, D	J											4390	D		
PCB-45	42.7	D				1000								534	D		
PCB-46	26.6	D												416	D		
PCB-47	137	D	J											2140	D		
PCB-48/75	46.1	D												523	D		
PCB-50	15.3	D		4.71	D, G		8.12	D, G	J	53.2	655	D, G					
PCB-51	32.1	D												436	D		
PCB-52/69	362	B, D	J											5220	D		
PCB-53	67.8	D	J											819	D		
PCB-56/60	189	B, D	J											2830	D	200000000000000000000000000000000000000	
PCB-61/70	345	D	l											5030	D		
PCB-63	15.3	D					3.23	D, G	J		614	D, G		202	Approximation of		101
PCB-67	9.10	D, G												101	D, G		
PCB-74	109	D												1720	D		
PCB-76/66	259	D	J											4020	D		
PCB-77	35.5	D												563	D		
PCB-79											396	D, G					
PCB-82	79.9	D	J	10.7	D	400	18.5	D	J	53.4	3340	D	J	1210	D		93.6

	PR1WWDUP-01C			PR1LDDUP-01C			PR1HDDUP-01C				PR1LPDUP-01C		PR1HPDUP-01C			
Analyte Identified	Whole Water ^g (pg/L)	LQ ^h	vq	LSM Dissolved [©] (pg/L)	LQ ^h	VQ	HSM Dissolved ⁸ (pg/L)	LQ ^h	VQ	% RPD	LSM Particulate ⁸ (pg/g)	ιQ ^h VQ	HSM Particulate ^g (pg/g)	LQh	VQ	% RPD
PCB-84/92	230	B, D	J								8300	DJ	3420	D		83.3
PCB-85/116	93.0	D		13.0	D		22.9	D	J	55.2	3830	DJ	1410			92.4
PCB-87/117/125	215	D	J								8330	L D	3150	D		90.2
PCB-88/91	77.8	D	J	12.5	D		19.7	D	J	44.7	3320	L D	1190	D		94.5
PCB-90/101	525	B, D	J				129	B, D	J		20400	B, D J	7520	D		92.3
PCB-95/98/102	390	B, D	J		100						15200	B, D J	5440	D		94.6
PCB-97	163	D	J								6290	DJ	2440	D		88.2
PCB-99	214	B, D	J				55.7	D	J		8040	D J	3330	D		82.8
PCB-105	209	D	J				43.9	D	J		7670	DJ	3100	D		84.9
PCB-106/118	503	B, D	J				105	B, D	J		19500	B, D J	7530	D		88.6
PCB-107/109	30.3	D		4.78	D, G		7.48	D, G	J	44.0	1570	D	564	D		94.3
PCB-108/112	30.2	D					7,35	D, G	J		1090	D, G	406	D		91.4
PCB-110	594	B, D	J				146	B, D	J		25500	B, D J	8940	D		96.2
PCB-111/115					2.74						669	D, G	165	D, G		121
PCB-114	11.7	D	J										187	D, G		
PCB-119	10.2	D, G									431	D, G	177	D, G		83.6
PCB-122													88.7	D, G		
PCB-123													185	D, G	J	
PCB-124	28.1	D					5.47	D, G	j		988	D, G	364	D		92.3
PCB-128/162	114	D	J	12.6	D		20.7	D	J	48.6	4220	DJ	1760	D		82.3
PCB-129	35.2	D	J				6.38	D, G	J		1500	D	475	D		104
PCB-130	47.4	D	J	4.36	D, G		7.63	D, G	J	54.5	1620	D	584	D		94.0
PCB-132/161	178	D	J								6780	DI	2750	D		84.6
PCB-133/142	16.0	D	J			Page Supplement	3.46	D, G	J		740	D, G	261	D		95.7
PCB-134/143	33.4	D	J	3.85	D, G		7.17	D, G	J	60.3	1270	D	481	D		90.1
PCB-135	75.7	D	J	13.3	D		20.7	D	J	43.5	4160	DJ	1310	D		104
PCB-136	75.7	D	J	9.13	D, G		17.6	D	J	63.4	3630	B, D J	1070	D		109
PCB-137	32.1	D	J	3.76	D, G		6.74	D, G	j	56.8	1350	D	406	D		107.5
PCB-138/163/164	674	B, D	J				114	B, D	J		25400	B, D J	9580	D		90.5
PCB-139/149	467	D	J	67.6	D		118	D	J	54.3	24100	B, D J	7260	D		107
PCB-141	151	D	J								4990	U O	1950	D		87.6
PCB-144	34.4	D					7.86	D, G	j		1530	D	402	D		116.8
PCB-146/165	77.3	D	J								2990	D J	1100	D		92.4
PCB-147											910	D, G				
PCB-151	138	B, D	J	17.8	D		31.3	D	J	55.0	6320	B, D J	1930	D		106
PCB-153	566	B, D	J				101	B, D	J		19900	B, D J	7790	D		87.5
PCB-156	72.1	D	J	7.31	D, G		10.8	a	J	38.5	2580	D J	1010	D		87.5
PCB-157	14.9	D	J	2.35	D, G		3.20	D, G	J	30.6	705	D, G	271	D		88.9
PCB-158/160	74.2	D	J								3110	D J	1100	D		95.5
PCB-167	31.3	D	J	3.89	D, G	100	5.18	D, G	J	28.4	1010	D, G	442	D		78.2
PCB-169																
PCB-170	231	D	J	15.6	D		29.4	D	J	61.3	7250	DJ	2900	D		85.7
PCB-171	61.8	D	J	4.47	D, G		7.89	D, G	J	55.3	1990	D I	677	D		98.5
PCB-172	46.5	D	J	3.86	D, G		6.40	D, G	J	49.5	1420	D	558	D		87.2

Analyte Identified	PR1WWDUP-01C Whole Water ^g (pg/L)	LQ ^h	VQ	PR1LDDUP-01C LSM Dissolved [®] (pg/L)	LQ ^h	VQ	PR1HDDUP-01C HSM Dissolved [®] (pg/L)	LQ ^h	۷Q	% RPD	PR1LPDUP-01C LSM Particulate ⁸ (pg/g)	LQ ^h	VQ	PR1HPDUP-01C HSM Particulate ^g (pg/g)	LQ ^h	VQ	% RPD
PCB-174	245	D	J	18.4	D		32.0	D	J	54.0	6750	l a		2740	D		84.5
PCB-175											359	D, G					
PCB-176	26.2	D	J	3.47	D, G	100	4.47	D, G	J	25.2	1020	D, G		308	D		107
PCB-177	136	D	J	10.6	D		18.6	D	J	54.8	4240	D J		1670	D		87.0
PCB-178	53.6	D	J	6.16	D, G		8.47	D, G	J	31.6	1930	DJ		666	D		97.4
PCB-179	97.0	D	J											1250	D		
PCB-180	540	B, D	J								15600	DJ		6430	D		83.3
PCB-182/187	302	D	J	28.8	D		44.2	D	J	42.2	11100	DJ		3730	D		99.4
PCB-183	131	D	J	12.2	D		19.9	D	j	48.0	4570	DJ		1690	D		92.0
PCB-184				3.63	D, G		4.98	D, G	J	31.4	610	D, G					
PCB-185	32.3	D	J								968	D, G		320	D		101
PCB-189														120	D, G		
PCB-190	47.6	D	J	3.29	D, G	200	6.12	D, G	J	60.1	1430	D		492	D		97.6
PCB-191	8.67	D, G	J								320	D, G					
PCB-193	25.4	D	J				3.4	D, G	J		699	D, G		331	D		71.5
PCB-194	137	D	J	6.79	D, G		15.3	D		77.0	3390	D J		1430	D		81.3
PCB-195	51.9	D	J	3,18	D, G		7.07	D, G		75.9	1230	D, G J		610	D		67.4
PCB-196/203	153	D	J	13.2	D		18.3	D	J	32.4	4910	DJ		1800	D		92.7
PCB-197											327	D, G					
PCB-199	157	D	J	11.5	D		17.9	D	J	43.5	5080	D J		1970	D		88.2
PCB-200	20.1	D	J											217	D		
PCB-201	22.0	D	J								685	D, G		234	D		98.2
PCB-202	36.3	D	J	3.15	D, G		6.06	D, G	J	63.2	1140	D, G J		430	D		90.4
PCB-206	105	D	J											1210	D		
PCB-207	11.2	D	J								251	D, G		167	D, G	J	40.2
PCB-208	30.0	D	J								945	D, G		412	D		78.6
PCB-209														1080	D		

a NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern FFS = focused fesability study HSM = high-solids mass

LSM = low-solids mass LQ = laboratory qualifier - See Attachment 1 for definitions PCB = polychlorinated biphenyl

pg/g = picograms per gram

pg/L = picograms per liter RPD = relative percent difference VQ = validation qualifier - See Attachment 2 for definitions

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPECs listed in the FFS: PCB -77, PCB -81, PCB -105, PCB -114, PCB -118, PCB -123, PCB -126, PCB -156, PCB -157, PCB -167, PCB -169, and PCB -189.

d At least 2

e Fewer than 17

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

g No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Appendix F	Н	X	dix	nen	An

Detailed Evaluation Sheets (Worksheet #11) - Aroclor PCBs

EVENT 1 ORIGINAL SAMPLE - AROCLOR PCBs PR1CSOCLY**-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Aroclor PCBs Sample Collection Techniques	Sample Col	lection Qualit	y ^a	Analytical Quality ^b	Identification	Identification of Target Analytes				
	Were specified sample aliquots obtained (r		Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Is at least 1 more COPC/COPEC ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?				
	Attempt 1	Attempt 2	Attempt 3							
Whole Water	Yes	Yes	NA	Yes	0	Yes	NA			
LSM dissolved plus LSM particulate	Yes	Yes	NA	Yes	0	Yes	NA			
HSM dissolved plus HSM particulate	Yes	Yes	NA	Yes	1	No	NA			
LSM dissolved	Yes	Yes	NA	Yes	0	No	No			
HSM dissolved	Yes	Yes	NA	Yes	0	140	No			
LSM particulate	Yes	Yes	NA	Yes	0	Yes	NA			
HSM particulate	No	Yes	NA	Yes	1	103	NA			

Positive Target Analyte Identification and Concentration Comparison[†]

Analyte I dentified	PR1CSOCLYWW-01B Whole Water ^g (µg/L)	LQ ^h	vq	PRICSOCLYLD-01B LSM Dissolved ^E (µg/L) LQ ^h VQ	PR1CSOCLYHD-01B HSM Dissolved ⁸ (µg/L) LQ ^h VQ	% RPD	PR1CSOCLYLP-01B LSM Particulate ^s (μg/kg) LQ ^h VQ	PR1CSOCLYHP-01B HSM Particulate ^g (µg/kg) LQ ^h VQ % RPD
Aroclor 1254								130 P J
Aroclor 1260								84 GP J

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern

FFS = focused fesability study HSM = high-solids mass LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCB = polychlorinated biphenyl RPD = relative percent recovery μg/L = micrograms per liter μg/Kg = micrograms per kilogram

VQ = validation qualifier - See Attachment 2 for definitions

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^cCOPCs/COPECs listed in the FFS: Aroclor 1016, Aroclor 1221, Aroclor 1232, Aroclor 1242, Aroclor 1248, Aroclor 1260, Aroclor 1262, and Arolcor 1268.

d At least 1 more

^e Fewer than 1

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

⁸ No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 1 FIELD DUPLICATE - AROCLOR PCBs PR1**DUP-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Aroclor PCBs Sample Collection Techniques	Sample Colle	ection Quali	: y *	Analytical Quality ^b	Identification	of Target Analy	tes
	Were specified sample aliquots obtained meeting all analytical needs? Attempt 1 Attempt 2 Atter			Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Is at least 1 more COPC/COPEC ^c identified in another sample type?	If no single sample type being compared was significantly ¹ different in the number of COPCs/COPECs ^c identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly ^c different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	Yes	Yes	NA	Yes	0	Yes	NA
LSM dissolved plus LSM particulate	Yes	Yes	NA	Yes	0	Yes	NA
HSM dissolved plus HSM particulate	Yes	Yes	NA	Yes	1	No	NA
LSM dissolved	Yes	Yes	NA	Yes	0	No	No
HSM dissolved	Yes	Yes	NA	Yes	0	110	No
LSM particulate	Yes	Yes	NA	Yes	0	Yes	NA
HSM particulate	No	Yes	NA	Yes	1	, es	NA

Positive Target Analyte Identification and Concentration Comparison

Analyte Identified	PR1WWDUP-01B Whole Water ^s (µg/L)	LQ ^h	VQ	PR1LDDUP-01B LSM Dissolved ² (µg/L)	LQ ^h VQ	PR1HDDUP-01B HSM Dissolved ^g (µg/L) LQ	' VQ	% RPD	PR1LPDUP-01B LSM Particulate ^g (µg/kg)	ra _r va	PR1HPDUP-01B HSM Particulate ^g (µg/kg)	
Aroclor 1254											160	M
Aroclor 1260				174					46		67	G M

a NNA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
EES = focused for ability ctudy

FFS = focused fesability study HSM = high-solids mass LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCB = polychlorinated biphenyl

RPD = relative percent recovery

μg/L = micrograms per liter

μg/Kg = micrograms per kilogram

VQ = validation qualifier - See Attachment 2 for definitions

 $^{^{\}mathrm{b}}$ Analytical quality is based upon the program 90% analytical completeness objectives.

COPCs/COPEcs listed in the FFS: Aroclor 1016, Aroclor 1221, Aroclor 1232, Aroclor 1242, Aroclor 1248, Aroclor 1260, Aroclor 1262, and Arolcor 1268.

d At least 1 more

e Fewer than 1

Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

⁸ No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 2 ORIGINAL SAMPLE - AROCLOR PCBs PR1CSOCLY**-02B

QAPP Worksheet #11-1

Project Quality Objectives/Systematic Planning Process Statements (Phase I)

Phase I Data Comparison Chart

Aroclor PCBs Sample Collection Techniques	Sample Colle	ection Qualit	yª	Analytical Quality ^b	Identification	Identification of Target Analytes					
	Were specified sample meeting all analytical			Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Is at least 1 more COPC/COPEC ^o identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?				
	Attempt 1	Attempt 2	Attempt 3								
Whole Water	No	Yes	NA	Yes	0	No	No				
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	0	No	No				
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	0	No	No				
LSM dissolved	No	Yes	NA	Yes	0	No	No				
HSM dissolved	No	Yes	NA	Yes	0	ING.	No				
LSM particulate	No	Yes	NA	Yes	0	No	No				
HSM particulate	No	Yes	NA	Yes	0	, NO	No				

Positive Target Analyte Identification and Concentration Comparison[†]

	PR1CSOCLYWW-02B Whole Water ^g (µg/L)	VQ	PR1CSOCLYLD-02B LSM Dissolved ⁶ (µg/L) LQ	ı va	PRICSOCLYHD-02B HSM Dissolved [®] (µg/L) LQ ^h VQ	% RPD	PR1CSOCLYLP-02B LSM Particulate [®] (µg/kg) LQ [®] VQ	PR1CSOCLYHP-02B HSM Particulate [#] (µg/kg) LQ ^h VQ % RPD
Aroclor 1254								47 G M

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern

LSM = low-solids mass

μg/L = micrograms per liter

COPECs = contaminants of potential ecological concern FFS = focused fesability study LQ = laboratory qualifier - See Attachment 1 for definitions PCB = polychlorinated biphenyl μg/Kg = micrograms per kilogram

HSM = high-solids mass

RPD = relative percent recovery

VQ = laboratory qualifier - See Attachment 2 for definitions

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

COPCs/COPECs listed in the FFS: Aroclor 1016, Aroclor 1221, Aroclor 1232, Aroclor 1242, Aroclor 1248, Aroclor 1260, Aroclor 1262, and Arolcor 1268.

^d At least 1 more

e Fewer than 1

¹ Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

^g No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

EVENT 2 FIELD DUPLICATE - AROCLOR PCBs PR1**DUP-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Aroclor PCBs Sample Collection Techniques	Sample Co	lection Quali	tyª	Analytical Quality ^b	ld entification o	ldentification of Target Analytes				
	Were specified samp meeting all analytica		otained	Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Is at least 1 more COPC/COPEC ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?			
	Attempt 1	Attempt 2	Attempt 3							
Whole Water	No	Yes	NA	Yes	0	Yes	NA			
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	0	Yes	NA			
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	1	No	NA			
LSM dissolved	No	Yes	NA	Yes	0	No	No			
HSM dissolved	No	Yes	NA	Yes	0	1,00	No			
LSM particulate	No	Yes	NA	Yes	0	Yes	NA			
HSM particulate	No	Yes	NA	Yes	1	1 '6'	NA			

Positive Target Analyte Identification and Concentration Comparison[†]

Analyte Identified	PR1WWDUP-02B Whole Water ^g (µg/L)	LQ ^h	VQ	PR1LDDUP-02B LSM Dissolved [®] (µg/L)	ιQ ^h	VQ	PR1HDDUP-02B HSM Dissolved [©] (µg/L)	LQ ^h VO	k % RPD	PR1LPDUP-02B LSM Particulate ^g (µg/kg)	LQ ^b VQ	PR1HPDUP-02B HSM Particulate ⁸ (µg/kg)	tq ^h vq 9	% RPD
Aroclor 1254												45	G M	
Aroclor 1260												22	GP J	

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern

FFS = focused fesability study

HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

PCB = polychlorinated biphenyl

RPD = relative percent recovery

μg/L = micrograms per liter

μg/Kg = micrograms per kilogram

VQ = laboratory qualifier - See Attachment 2 for definitions

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

COPCs/COPEs listed in the FFS: Aroclor 1016, Aroclor 1221, Aroclor 1232, Aroclor 1242, Aroclor 1248, Aroclor 1260, Aroclor 1262, and Arolcor 1268.

^d At least 1 more

e Fewer than 1

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

⁸ No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.



EVENT 1 ORIGINAL SAMPLE - ORGANOCHLORINE PESTICIDES PR1CSOCLY**-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Organochlorine Pesticides Sample Collection Techniques	Sample Coll	ection Quali	t ∛	Analytical Quality ^b	Identification of	Target Analytes	
	Were specified sample meeting all analytical r		ined	Are fewer than 4 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Is at least 1 more COPC/COPEC [©] identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	Yes	Yes	NA	Yes	3	No	No
LSM dissolved plus LSM particulate	Yes	Yes	NA	Yes	3	No	No
HSM dissolved plus HSM particulate	Yes	Yes	NA	No (4) ^f	NA	NA	NA
LSM dissolved	Yes	Yes	NA	Yes	3	No	Yes
HSM dissolved	Yes	Yes	NA	Yes	3	NO	Yes
LSM particulate	Yes	Yes	NA	Yes	2	Yes	NA
HSM particulate	No	Yes	NA	No (4) ^f	NA.	NA	NA NA

Positive Target Analyte Identification and Concentration Comparison^g

Analyte Identified	PR1CSOCLYWW-01B Whole Water ^h (pg/L)	LQ ⁱ	VQ	PR1CSOCLYLD-01B LSM Dissolved (pg/L)	LQ ^l	VQ	PR1CSOCLYHD-01B HSM Dissolved ^h (pg/L)	ια'	VQ	% RPD	PR1CSOCLYLP-01B LSM Particulate ^h (pg/g)	LQ ^l	VQ	PR1CSOCLYHP-01B HSM Particulate (pg/g)	ια'	vq	% RPD
alpha-BHC				25.8			400				112						
Lindane (gamma-BHC)	313		J	262			291		J	10.5	455			294		J	43.0
beta-BHC	136		j	110			131		J	17.4				71.9	G	J	
Heptachlor	151			70.9	G		130		J	58.8	1300	DG	j	138	G	J	162
Aldrin	82.3		J	36.8		J	65		J	55.4	772		J				
Oxychlordane	46.9		J				44.9		J		646		J				
cis-Heptachlor Epoxide	371		J	210			320		j	41.5	2600		J	555		J	130
trans-Chlordane (gamma)	2020		j	865		J	1870		J	73.5	202000		J	3930		J	192
trans-Nonachlor	1190		J	422		j	774		J	58.9	8890		J	2780		J	105
cis-Chlordane (alpha)	2270	D	J	1120		J	1870		J	50.2	17800		j	5320		j	108
Endosulfan I (alpha)	112	G	J	70.3	U	J	82.5	G	J	16.0							
4,4'-DDE														7840		J	
Dieldrin	2450	BD	J	1160	В	J	2390	BD	j	69.3				3680		j	
Endrin							28.6	G	J								
cis-Nonachlor	257		J	117		J	252		J	73.2	1820		J	538		J	109
Endosulfan II (beta)							85.4	G	J								
4,4'-DDD														29200	E	J	
Endosulfan Sulfate							101	G	J								
4,4'-Methoxychlor	480		J	239		j	380		1	45.6	3980	DG		ND	U	R ^j	
Mirex							16.5		J					ND	U	R ^j	
Endrin Aldehyde														ND	RINGSONNIU QUINOS	R ^j	
Endrin Ketone	97.1	G	J	85	В	J	64.6	G	J	27.3				ND	U	R ^j	

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study
HSM = high-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions pg/g = picograms per gram pg/L = picograms per liter R = rejected data result

RPD = relative percent difference VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

^b Analytical quality is based upon the program 90% analytical completeness objectives.

 $^{^{\}circ}$ COPCs/COPECs listed in the FFS: cis-Chlordane(alpha), trans-Chlordane(gamma), Dieldrin, 4,4'-DDE, 4,4'-DDD, and 4,4'DDT.

^d At least 1 more

e Fewer than 4

^fValues in parentheses indicate the total number of rejected results

⁸ Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

h No rejected data.

A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitationely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

PRICSOCLYHP-01B- All data results rejected due to low labeled analog standard recovery. Sample not used during sample collection technique evaluation.

EVENT 1 FIELD DUPLICATE - ORGANOCHLORINE PESTICIDES PR1**DUP-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Organochlorine Pesticides Sample Collection Techniques	Sample Co	llection Qua	lity ^a	Analytical Quality ^b	ldentification	of Target Analyt	es
	Were specified san meeting all analyti		s obtained	Are fewer than 4 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Is at least 1 more COPC/COPEC ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	Yes	Yes	NA	Yes	3	Yes	NA
LSM dissolved plus LSM particulate	Yes	Yes	NA	Yes	3	Yes	NA
HSM dissolved plus HSM particulate	Yes	Yes	NA	Yes (2) ^f	5	No	NA
LSM dissolved	Yes	Yes	NA	Yes	3	No	Yes
HSM dissolved	Yes	Yes	NA	Yes	3	NO NO	Yes
LSM particulate	Yes	Yes	NA	Yes	3	Yes	NA
HSM particulate	No	Yes	NA	Yes (2) ^f	5	163	NA

Positive Target Analyte Identification and Concentration Comparison⁸

Analyte Identified	PR1WWDUP-01B Whole Water ^h (pg/L)	LQ ⁱ	VQ	PR1LDDUP-01B LSM Dissolved ^h (pg/L)	ια ^ί	VQ	PR1HDDUP-01B HSM Dissolved ^h (pg/L)	ια'	۷Q	% RPD	PR1LPDUP-01B LSM Particulate ¹ (pg/g)	LQ ^I	VQ	PR1HPDUP-01B HSM Particulate (pg/g)	Γά	VQ	% RPD
alpha-BHC	26.5		J														
Lindane (gamma-BHC)	311		J	286		J	290		J	1.4	617		J	319		J	63.7
beta-BHC	127		J	124		J	128		J	3.2	520		J	268		J	64.0
delta-BHC							6.46	G	J								
Heptachlor	143		J				129		J		1290		J	470	G	J	93.2
Aldrin	88.7		J	40.5		J	55.8		J	31.8							
Oxychlordane	60.6		J											476		J	
cis-Heptachlor Epoxide	376		J	211		J	335		J	45.4	2770		J	1690		J	48.4
trans-Chlordane (gamma)	1880		J	1020	D		1590		J	43.7	22100		J	10900		Į	67.9
trans-Nonachlor	1070		J	605		j	935		J	42.9	10800		J	7350		J	38.0
cis-Chlordane (alpha)	2440	D	J	1120		J	1830	D		48.1	21800		J	15200	E	J	35.7
Endosulfan I (alpha)	121	G	J	10.46			117	G	J		1050	G	J				
4,4'-DDE														23000		J	172
Dieldrin	2610	BD	J	1240	В	J	2290	BD	J	59.5	18000		J	9470		J	62.1
cis-Nonachlor	290		J	Number of the Control	il.		100				2480		J	2750		J	10.3
Endosulfan II (beta)							112	G	J								
4,4'-DDD														102000	E	J	
Endosulfan Sulfate							112	G	J								
4,4'-Methoxychlor	523		J	257	DG	J	375		J	37.3	3410		J	ND	U	R ^J	
Endrin Aldehyde														ND	U	R	
Endrin Ketone							83.1		J		447.483						

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern FFS = focused fesability study HSM = high-solids mass LQ = laboratory qualifier - See Attachment 1 for definitions pg/g = picograms per gram pg/L = picograms per liter R = rejected data result

RPD = relative percent difference VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPECs listed in the FFS: cis-Chlordane(alpha), trans-Chlordane(gamma), Dieldrin, 4,4'-DDE, 4,4'-DDD, and 4,4'DDT.

d At least 1 more

^e Fewer than 4

^f Values in parentheses indicate the total number of rejected results

⁸ Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

^h No rejected data.

¹ A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

PRIHPDUP-01B All data results rejected due to low labeled analog standard recovery. Evaluation was not impacted based on rejected result.

EVENT 2 ORIGINAL SAMPLE - ORGANOCHLORINE PESTICIDES PR1CSOCLY**-02B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

Organochlorine Pesticides Sample Collection Techniques	Sample Coll	ection Qual	ity ^a	Analytical Quality ^b	Identification of	Target Analyte	s
	Were specified samp meeting all analytica		btained		Number of COPCs/COPECs ^c listed in the FFS identified?	Is at least 1 more COPC/COPEC ^c identified in another sample type?	If no single sample type being compared was significantly ^d different in the number of COPCs/COPECs ^c identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly ^e different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	No	Yes	NA	Yes	3	Yes	NA
LSM dissolved plus LSM particulate	No	Yes	NA	Yes (1) ^f	3	Yes	NA
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	4	No	NA
LSM dissolved	No	Yes	NA	Yes	3	No	No
HSM dissolved	No	Yes	NA	Yes	3	140	No
LSM particulate	No	Yes	NA	Yes (1) ^f	3	Yes	NA
HSM particulate	No	Yes	NΑ	Yes	4	i es	NA

Positive Target Analyte Identification and Concentration Comparison⁸

Analyte Identified	PR1CSOCLYWW-02B Whole Water ^h (pg/L)	LQ ¹	VQ	PR1CSOCLYLD-02B LSM Dissolved ^h (pg/L)	ΙQ	VQ	PR1CSOCLYHD-02B HSM Dissolved ^h (pg/L)	Ġ	VQ	% RPD	PR1CSOCLYLP-02B LSM Particulate (pg/g)	ιQ	VQ	PR1CSOCLYHP-02B HSM Particulate ^h (pg/g)	ια ^ί	VQ	% RPD
Hexachlorobenzene														2670	D	J	
alpha-BHC	70.1			66.9			60.3		J	10.4				102	D	J	
Lindane (gamma-BHC)	146			147			153		J	4.0				342	D	J	
beta-BHC	23													223	D	j	
Heptachlor	43.9	G					43.2	G	J					680	D	J	
Aldrin											1290	G					
Oxychlordane	33.4		J								2710			554	D	J	132
cis-Heptachlor Epoxide	128		···	65.0			112		J	53.1	6060			1590	D	J	117
trans-Chlordane (gamma)	674			210		J	513		j	83.8	62600			10000	D	М	145
trans-Nonachlor	439			123		j	311		J	86.6	39500		J	8080	D	J	132
cis-Chlordane (alpha)	661		j	218		J	591		J	92.2	67500		J	13500	D	J	133
Endosulfan I (alpha)	64.4	G					53.7	G	J		2960	G	J				
4,4'-DDE														21100	D	J	
Dieldrin	421			220			480		J	74.3	27300		J	5050	D	J	138
cis-Nonachlor	113			33.6			80.6		J	82.3	11800		j	2320	D	JH	134
Endosulfan II (beta)	633		J	64.9	G		93.5	G	J	36.1							
Endosulfan Sulfate				45.0	G												
4,4'-Methoxychlor	170	G	J	67.0	G		120		J	56.7	11500	G	J				
Endrin Aldehyde		_									ND	U	R ^j				
Mirex				2.29	G	J			J		1090	G	J				

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study

FFS = focused fesability study HSM = high-solids mass LQ = laboratory qualifier - See Attachment 1 for definitions pg/g = picograms per gram pg/L = picograms per liter R = rejected data result

RPD = relative percent difference $VQ = laboratory\ qualifier - See\ Attachment\ 2\ for\ definitions$ % = percent

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPECs listed in the FFS: cis-Chlordane(alpha), trans-Chlordane(gamma), Dieldrin, 4,4'-DDE, 4,4'-DDD, and 4,4'DDT.

d At least 1 more

e Fewer than 4

^f Values in parentheses indicate the total number of rejected results

g Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

h No rejected data.

i A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

¹ PR1CSOCLYLP-02B Data result rejected due to low labeled analog standard recovery. Evaluation was not impacted based on rejected result.

EVENT 2 FIELD DUPLICATE - ORGANOCHLORINE PESTICIDES PR1**DUP-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Organochlorine Pesticides Sample Collection Techniques	Sample Col	lection Qual	ity ^a	Analytical Quality ^b	Identification	of Target Analyt	es
	Were specified sam meeting all analytic		obtained	Are fewer than 4 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	ls at least 1 more COPC/COPEC ^c identified in another sample type?	If no single sample type being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	No	Yes	NA	Yes	3	No	No
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	3	No	No
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	3	No	No
LSM dissolved	No	Yes	NA	Yes	3	No	No
HSM dissolved	No	Yes	NA	Yes	3	140	No
LSM particulate	No	Yes	NA	Yes	3	No	No
HSM particulate	No	Yes	NA	Yes	3	No	No

Positive Target Analyte Identification and Concentration Comparison[†]

Analyte Identified	PR1WWDUP-02B Whole Water ^g (pg/L)	LQ ^h	VQ	PR1LDDUP-02B LSM Dissolved ^g (pg/L)	LQ ^h	VQ	PR1HDDUP-02B HSM Dissolved ⁵ (pg/L)	ιQ ^h	VQ	% RPD	PR1LPDUP-02B LSM Particulate ⁶ (pg/g)	ΓŌμ	VQ	PR1HPDUP-02B HSM Particulate ⁶ (pg/g)	LQ ^b	VQ	% RPD
alpha-BHC	72.7			63.5			63.2			0.47				82.7	DG	M	
Lindane (gamma-BHC)	147			134			150			11.3				203	DG	J	
beta-BHC	30.6						20							231	DG	M	
Heptachlor							41.2	G			2890	G					
Aldrin											997	G		264	DG	J	116
Oxychlordane	44.6		J								2110			460	DG	M	128
cis-Heptachlor Epoxide	137		J	56.2			119			71.7	4870			1530	D	M	104
trans-Chlordane (gamma)	648			204		J	540			90.3	49800			9350	D	M	137
trans-Nonachlor	421		J	120		J	320		J	90.9	27400		J	7790	D	M	111
cis-Chlordane (alpha)	665		J	200		J	622		J	103	55600		J	13600	D	M	121
Endosulfan I (alpha)				41.6	G	J	52.2	G	J	22.6	1850	G	J	502	DG	J	115
Dieldrin	449		J	214			456		J	72.2	18200		J	5550	D	J	107
cis-Nonachlor	115		J	33.7			81.8		J	83.3	7820		J	2740	D	J	96.2
Endosulfan II (beta)	711		J				80.6	G	J								
Endosulfan Sulfate	117	G	J	47.9	G												
4,4'-Methoxychlor	174		J	62.7	G		107	G	J	52.2	6960	G	J				
Mirex	13.8	G	J														
Endrin Ketone				10.9	G												

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study

HSM = high-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions pg/g = picograms per gram pg/L = picograms per liter RPD = relative percent difference VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPECs listed in the FFS: cis-Chlordane(alpha), trans-Chlordane(gamma), Dieldrin, 4,4'-DDE, 4,4'-DDD, and 4,4'DDT.

^d At least 1 more

e Fewer than 4

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full Target Analyte List. Additional pages may be necessary.

^g No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitationally less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Appendix J

Detailed Evaluation Sheets (Worksheet #11) - SVOCs

EVENT 1 ORIGINAL SAMPLE - SEMIVOLATILES PR1CSOCLY**-01B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

SVOC Sample Collection Techniques	Sample Coll	ection Quality	î'	Analytical Quality	Identification of Target	Analytes
	Were specified sample a all analytical needs?	liquots obtain	ed meeting	Are fewer than 6 results "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least five more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	Yes	Yes	NA	Yes	4	NA
LSM dissolved plus LSM particulate	Yes	Yes	NA	No (9) ^c	NA	NA
HSM dissolved plus HSM particulate	Yes	Yes	NA	No (8) ^c	NA	NA
LSM dissolved	Yes	Yes	NA	Yes (1) ^c	3	NA
HSM dissolved	Yes	Yes	NA	No (8) ^c	NA NA	NA
LSM particulate	Yes	Yes	NA	No (9) ^c	NA	NA
HSM particulate	No	Yes	NA	Yes (1)°	2	NA

Positive Target Analyte Identification and Concentration Comparison d

Analyte Identified	PR1CSOCLYWW-01B Whole Water ^e (μg/L)	LQ ^f	VQ	PR1CSOCLYLD-01B LSM Dissolved (μg/L)	ιq'	VQ	PR1CSOCLYHD-01B HSM Dissolved (µg/L)	ιQ ^f	VQ	% RPD	PR1CSOCLYLP-01B LSM Particulate (µg/kg)	LQ'	VQ	PR1CSOCLYHP-01B HSM Particulate (μg/kg)	ιq'	VQ	% RPD
Phenoi				2.4			1.7		J	34.1							
4-Methylphenol	0.80	GD		9.3		J	5.4		J	53.1				5100	GD	М	
Diethylphthalate	3.1	D															
Di-n-butylphthalate	2.2	DB		0.70	GB		2.7		J	118	4100	GD	J	13000	DB	M	104
Butylbenzylphthalate							2.8	В	3 J								
Bis(2-ethylhexyl)phthalate	5.3	DB					29	EE	} J								
Di-n-octylphthalate				ND	U	R ⁸	ND	ι	ı R ^h		ND	U	R ^h	ND	U	R ⁸	
4,6-Dinitro-2-methylphenol							ND	ι	J R ^h		ND	U	R ^h				
N-Nitrosodiphenylamine							ND	L	J R ^h		ND	U	R ^h				
4-Bromophenyl-phenylether							ND	L	J R ^h		ND	U	R ^h				
Hexachlorobenzene							ND	L	J R ^b		ND	U	R ^h				
Atrazine							ND	ι	J R ^h		ND	U	R ^h				
Pentachlorophenol							ND	ι	J R ^h		ND	U	R ^h				
Carbazole			·				ND	ι	J R ^h		ND	U	R ^h				·
3,3'-Dichlorobenzidine							100				ND	U	R ^h				

There are no COPC/COPECs in the target list for SVOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions R = rejected data result RPD = relative percent difference SVOC = semivolatile organic compound

 μ g/L = micrograms per liter μ g/kg = micrograms per kilograms VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

a Na" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

 $^{^{\}circ}$ Values in paretheses indicate the total number of rejected results.

d Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^e No rejected data.

f A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

⁸ PR1CSOCLYLD-01B and PR1CSOCLYHP-01B data results rejected due to low internal standard recovery. Sample collection technique evaluation not impacted based on rejected results.

h PR1CSOCLYHD-01B and PR1CSOCLYLP-01B data results rejected due to low internal standard recovery. Samples not used during sample collection technique evaluation.

EVENT 1 FIELD DUPLICATE - SEMIVOLATILES PR1**DUP-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

SVOC Sample Collection Techniques	Sample C	ollection Qual	itvª	Analytical Quality b	Identification of Targ	et Analytes
	Were specified sam meeting all analytic	al needs?		Are fewer than 6 results "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have a least five more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3		<u> </u>	
Whole Water	Yes	Yes	NA	Yes	4	No
LSM dissolved plus LSM particulate	Yes	Yes	NA	Yes (1) ^c	4	No
HSM dissolved plus HSM particulate	Yes	Yes	NA	No (8) ^c	NA	NA
LSM dissolved	Yes	Yes	NA	Yes	4	NA
HSM dissolved	Yes	Yes	NA	No (8) ^c	NA	NA
LSM particulate	Yes	Yes	NA	Yes (1) ^c	2	No
HSM particulate	No	Yes	NA	Yes (1) ^c	4	No

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1WWDUP-01B Whole Water ^e (µg/L)	LQ ^f	VQ	PR1LDDUP-01B LSM Dissolved ^e (μg/L)	ַב <u>ל</u>	VQ	PR1HDDUP-01B HSM Dissolved (µg/L)	ra,	VQ	% RPD	PR1LPDUP-018 LSM Particulate (µg/kg)	LQ ^f	VQ	PR1HPDUP-01B HSM Particulate (µg/kg)	ιqʻ	VQ	% RPD
Phenol	2.1	GD					2.0	GD									
Acetophenone				0.30	G		100000000000000000000000000000000000000										
4-Methylphenol							8.6	D	J					4000		M	
Diethylphthalate	3.7	D		3.7		J	3,4	D	J	8.45	2200	G					
Di-n-butylphthalate	3.0	GDB		1,1	В		2.1	GD	j	62.5	5900	G		4200	В	М	33.7
Butylbenzylphthalate				1.7	В									37000	EB	J	
Bis (2-ethylhexyl) phthalate	8.3	DB												25000	EB	ı	
Di-n-octylphthalate							ND	U	R ⁸		ND	U	R ^h	ND	U	R ^h	
4,6-Dinitro-2-methylphenol							ND	U	R ^g								
N-Nitrosodiphenylamine							ND	U	R ⁸								
4-Bromophenyl-phenyl ether							ND	U	R ^g								
Hexachlorobenzene							ND	U	R ^g								
Atrazine			•				ND	U	R ^g								
Pentachlorophenol			•				ND	U	R ^g								
Carbazole							ND	U	R ^g								

There are no COPC/COPECs in the target list for SVOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions
R = rejected data result
RPD = relative percent difference
SVOC = semivolatile organic compound

 $\mu g/L$ = micrograms per liter $\mu g/kg$ = micrograms per kilograms VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

a Na" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

 $^{^{\}rm c}$ Values in paretheses indicate the total number of rejected results.

d Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^e No rejected data.

A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

⁸ PR1HDDUP-01B data results rejected due to low internal standard recovery. Sample not used during sample collection technique evaluation.

h PR1LPDUP-01B and PR1HPDUP-01B data results rejected due to low internal standard recovery. Sample collection technique evaluation not impacted based on rejected results.

EVENT 2 ORIGINAL SAMPLE - SEMIVOLATILES PR1CSOCLY**-02B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

SVOC Sample Collection Techniques	Sample Coll	ection Qualit	y ^a	Analytical Quality ^b	Identification of Target	Analytes
	Were specified sample all analytical needs?	aliquots obtai	ned meeting	Are fewer than 6 results "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least five more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	No	Yes	NA	Yes	4	No
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	5	No
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	10	Yes
LSM dissolved	No	Yes	NA	Yes	4	No
HSM dissolved	No	Yes	NA	Yes	5	No
LSM particulate	No	Yes	NA	Yes	1	No
HSM particulate	No	Yes	NA	Yes	8	Yes

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1CSOCLYWW-02B Whole Water ^d (µg/L)	LQ*	VQ	PR1CSOCLYLD-02B LSM Dissolved ^d (µg/L)	LQ"		PR1CSOCLYHD-02B HSM Dissolved ^d (µg/L)	LQ*	VQ	% RPD	PR1CSOCLYLP-02B LSM Particulate ^d (μg/kg)	LQ°	VQ	PR1CSOCLYHP-02B HSM Particulate ^d (μg/kg)	LQ" VC	% RPD
Phenol				0.27	G	j	0.29	G		7.14						
Acetophenone	0.17	G		0.16	G		0.17	G		6.06						
4-Methylphenol														120	GJ	
Dibenzofuran														48	GМ	
Diethylphthalate	1.3			1.3			1.3			0.00				35	G M	
Carbazole														300	GМ	
Di-n-butylphthalate	0.22	G		0.24	G		0.28	G		15.4				320	GМ	
Butylbenzylphthalate														1200	М	
Bis(2-ethylhexyl)phthalate	2.5	·	•	100			2.1				240000)		12000	J J	181
Di-n-octylphthalate		·	•											2000	J	

There are no COPC/COPECs in the target list for SVOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

RPD = relative percent difference SVOC = semivolatile organic compound

μg/L = micrograms per liter

μg/kg = micrograms per kilograms

VQ = laboratory qualifier - See Attachment 2 for definitions

a NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

d No rejected data.

e A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 2 FIELD DUPLICATE - SEMIVOLATILES PR1**DUP-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

SVOC Sample Collection Techniques	Sample Co	llection Qua	lity ^a	Analytical Quality ^b	Identification of Targe	t Analytes
	Were specified san meeting all analyti		obtained	Are fewer than 6 results "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least five more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	No	Yes	NA	Yes	4	No
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	5	No
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	8	No
LSM dissolved	No	Yes	NA	Yes	4	No
HSM dissolved	No	Yes	NA	Yes	4	No
LSM particulate	No	Yes	NA	Yes	1	No
HSM particulate	No	Yes	NA	Yes	6	Yes

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1WWDUP-02B Whole Water ^d (µg/L)	LQ°	VQ	PR1LDDUP-02B LSM Dissolved ^d (µg/L)	LQ*	VQ	PR1HDDUP-02B HSM Dissolved ^d (µg/L)	LQ°	VQ	% RPD	PR1LPDUP-02B LSM Particulate ^d (µg/kg)	LQ ^e	VQ	PR1HPDUP-02B HSM Particulate ^d (μg/kg)	ια°	VQ	% RPD
Phenol	0.18	G	J	0.32	G		0.28	G		13.3							
Acetophenone				0.14	G												
4-Methylphenol														66	G	J	
Dibenzofuran				19										42	G	М	
Diethylphthalate	1.6			1.1			1.1			0.00							
Carbazole														130	G	М	
Di-n-butylphthalate	0.32	G		0.20	G		0.28	G		33.3				250	G	M	
Butylbenzylphthalate														1400		М	
Bis (2-ethylhexyl) phthalate	3.0	·					2.3				180000			11000	D	J	177

There are no COPC/COPECs in the target list for SVOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions RPD = relative percent difference SVOC = semivolatile organic compound $\mu g/L = micrograms \ per \ liter$

µg/kg = micrograms per kilograms VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^d No rejected data.

^e A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Λ							14	,
Α	р	р	eı	nc	11	X	n	١

Detailed Evaluation Sheets (Worksheet #11) – SVOCs SIM

EVENT 1 ORIGINAL SAMPLE- SEMIVOLATILES-SIM PR1CSOCLY**-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

SVOC SIM	I			ı	1		
Sample Collection Techniques	Sample Co	llection Quality	ř	Analytical Quality	Identification	of Target Analyt	es
	Were specified sample all analytical needs?	aliquots obtair	ned meeting	Are fewer than 3 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/COPECs ^c identified in another sample type?	If no single sample types being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3			<u> </u>	
Whole Water	Yes	Yes	NΑ	Yes	12	Yes	NA
LSM dissolved plus LSM particulate	Yes	Yes	NA	Yes	10	Yes	NA
HSM dissolved plus HSM particulate	Yes	Yes	NΑ	Yes	16	No	NA
LSM dissolved	Yes	Yes	NA	Yes	4	Yes	NA
HSM dissolved	Yes	Yes	NA	Yes	7		NA
LSM particulate	Yes	Yes	NA	Yes	6	Yes	NA NA
HSM particulate	No	Yes	NA	Yes	14		NA

Positive Target Analyte Identification and Concentration Comparison

							PR1CSOCLYHD-01B				PR1CSOCLYLP-01B			PR1CSOCLYHP-01B			
Analyte Identified	PR1CSOCLYWW-01B Whole Water ^g (µg/L)	LQ ^h	VQ	PR1CSOCLYLD-01B LSM Dissolved ⁸ (µg/L)	10 ^h	vq	HSM Dissolved ^g (µg/L)	LQ	VQ	% RPD	LSM Particulate ^s (µg/kg)	LQ ^h	vq	HSM Particulate ^ε (μg/kg)	LQ ^h	VQ	% RPD
Naphthalene	0.26	DB .		0.34			0.24	DB		34.5	[10" 104]		1	(46/16)		, , ,	7011112
								10000						440			
2-Methylnaphthalene	0.32	DB .		0.41	DB		0.34	DB		18.7			1	110	DB	J	
Acenaphthene	0.023	D.		0.022	D		0.019	D	1	14.6							
Fluorene	0.031	DB .		0.021	D		0.025	DB		17.4				75			
Phenanthrene	0.11	DB .					0.076	DB	IJ					710	DB		
Anthracene	0.022	DB .	J	10.4%										120	D	0200 00000	
Fluoranthene	0.15	DB .	J				0.054	DB			870			1900	DB	0.000	74.4
Pyrene	0.15	DB .	J				0.083	DB	J		930	DB	1	1000	DB		7.25
Benzo (a) anthracene							200							780	D	J	
Chrysene				1000000									ļ	920	D	J	
Benzo(b)fluoranthene	0.050	DB .	J								630	D		890	D	J	34.2
Benzo(k)fluoranthene	0.049	DB .	J								500	ס		730	D	J	37.4
Benzo(a) pyrene	0.038	DB .	J								450	D		750	D	J	50.0
Indeno(1,2,3-cd)pyrene														400	D	J	
Dibenzo (a,h) an thracene														120	D	j	
Benzo(g,h,i)perylene	0.022	DB .	J								310	D		410	D	J	27.8
1-Methylnaphthalene	0.22	DB .	J	0.28	D		0.23	DB	J	19.6				68	D	J	
Benzo[e]pyrene	0.036	DB .	J								420	D		640	D	J	41.5
Perylene														200	D	j	
3,6-Dimethylphenanthrene				100										54	D	j	
1-Methylanthracene	0.049	DB .	j	0.031	D	j	0.050	DB	J	46.9	620	D	J	260	D	J	81.8
1-Methylfluoranthene											310	D		180	D	J	53.1
1-Methylpyrene				1										87	D	j	
2,6-Dimethylnaphthalene	0.16	DB .	J	0.10	D		0.14	DB	ı J	33.3	480	D		150	D	J	104.8
2,3,5-Trimethylnaphthalene	0.092	DB .	J	0.054	D		0.070	DB		25.8	580	GD		120	D	j	131.4
1,1'-Biphenyl	0.022	DB .	J				0.019	DB	J								
1-Methylphenanthrene	0.084	DB .	J	0.037	D		0.069	DB		60.4				190		j	
Dibenzothiophene	0.029	DB .	j				0.026							51		j	

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study
HSM = high-solids mass
LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions RPD = relative percent difference SIM = selective ion monitoring SVOC = semivolatile organic compound µg/L = micrograms per liter

µg/kg = micrograms per kilograms
VQ = laboratory qualifier - See Attachment 2 for definitions
% = percent

^b Analytical quality is based upon the program 90% analytical completeness objectives.

[°] COPCs/COPECs listed in the FFS: Naphthalene, Fluorene, Pyrene, Benzo(k)fluoranthene, Benzo(g,h,i)perylene, 2-methylnaphthalene, Phenanthrene, Benzo(a)anthracene,

Benzo(a)pyrene, Acenaphthylene, Anthracene, Chrysene, Indeno(1,2,3-cd)pyrene, Acenaphthene, Fluoranthene, Benzo(b)fluoranthene, and Dibenzo(a,h)anthracene.

^d At least 2 more

e Fewer than 3

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^g No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitationally less certain than those not associated with a "G" qualifier are quantitationally less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 1 FIELD DUPLICATE - SEMIVOLATILES-SIM PR1**DUP-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

SVOC SIM Sample Collection Techniques	Sample C	ollection Qua	lity ^a	Analytical Quality ^b	Identification	on of Target Analyte	es
	Were specified san meeting all analyti		obtained	Are fewer than 3 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/COPECs ^c identified in another sample type?	If no single sample types being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	Yes	Yes	NA	Yes	9	Yes	NA
LSM dissolved plus LSM particulate	Yes	Yes	NA	Yes	11	Yes	NA
HSM dissolved plus HSM particulate	Yes	Yes	NA	Yes	14	No	NA
LSM dissolved	Yes	Yes	NA	Yes	4	No	Yes
HSM dissolved	Yes	Yes	NA	Yes	5	IVO	Yes
LSM particulate	Yes	Yes	NA	Yes	7	Yes	NA
HSM particulate	No	Yes	NA	Yes	12	163	NA

Positive Target Analyte Identification and Concentration Comparison

Analyte Identified	PR1WWDUP-01B Whole Water ^s (μg/L)	LQ ^h	VQ	PR1LDDUP-01B LSM Dissolved ^g (μg/L)	LQ ^h	VQ	PR1HDDUP-01B HSM Dissolved ^g (µg/L)	LQ ^b	VQ	% RPD	PR1LPDUP-01B LSM Particulate ⁸ (μg/kg)	LQ ^h	VQ	PR1HPDUP-01B HSM Particulate ^g (µg/kg)	LQ ^h	VQ	% RPD
Naphthalene	0.30	BD	J	0.37	DB		0.23	DB	J	46.7							
2-Methylnaphthalene	0.40	BD	J	0.44	D		0.31	DB	j	34.7				71	DB	J	
Acenaphthene				0.020	D												
Fluorene	0.028	BD	J	0.022	D		0.020	DB	j								
Phenanthrene	0.097	BD	J				0.063	DB	J					300	DB	J	
Fluoranthene	0.12	BD	J								1600	DB	J	770	DB	J	70.0
Pyrene	0.14	BD	J				0.069	DB	J		1000	DB	J	680	DB	J	38.1
Benzo(a)anthracene				A CONTRACTOR OF THE PARTY OF TH										310	D	j	
Chrysene														410	D	j	
Benzo(b)fluoranthene	0.042	BD	J	Section 200							880	D		390	D	j	77.2
Benzo(k) fluoranthene	0.043	BD	J								720	D		290	D	J	85.1
Benzo(a)pyrene	0.033	BD	J				10 mg				540	D		280	D	J	63.4
Indeno(1,2,3-cd)pyrene											300	D		180	D	J	50.0
Dibenzo(a,h)anthracene														66	D	j	
Benzo(g,h,i)perylene											340	D		220	D	J	42.9
1-Methylnaphthalene	0.26	BD	J	0.31	D		0.21	DB	j	38.5							
Benzo[e]pyrene	0.029	BD	J								550	D		270	D	j	68.3
Perylene				50										77	D	j	
3,6-Dimethylphenanthrene											330	D					
1-Methylanthracene	0.040	BD	J	0.030	a	J	0.043	DB	J	35.6	630	D	J	91	D	J	150
1-Methylfluoranthene											320	D		110	D	J	97.7
2,6-Dimethylnaphthalene	0.15	BD	J	0.10	D		0.12	DB	J	18.2	450	D		100	D	J	127
2,3,5-Trimethylnaphthalene	0.083	BD	J	0.054	D		0.074	DB	J	31.3	700	D		76	D	J	161
1-Methylphenanthrene	0.082	BD	j	0.037	D		0.061	DB	J	49.0							
Dibenzothiophene	0.028	BD]				0.025	DB	j								

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study
HSM = high-solids mass
LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions RPD = relative percent difference $SIM = selective ion monitoring \\ SVOC = semivolatile organic compound \\ \mu g/L = micrograms per liter$

 $\mu g/kg$ = micrograms per kilograms VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPECs listed in the FFS: Naphthalene, Fluorene, Pyrene, Benzo(k)fluoranthene, Benzo(g,h,i)perylene, 2-methylnaphthalene, Phenanthrene, Benzo(a)anthracene, Benzo(a)pyrene, Acenaphthylene, Anthracene, Chrysene, Indeno(1,2,3-cd)pyrene, Acenaphthene, Fluoranthene, Benzo(b)fluoranthene, and Dibenzo(a,h)anthracene.

^d At least 2 more

e Fewer than 3

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

⁸ No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 2 ORIGINAL SAMPLE - SEMIVOLATILES-SIM PR1CSOCLY**-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

SVOC SIM Sample Collection Techniques	Sample C	ollection Qualit	ŧγ³	Analytical Quality	Identification of Target Analytes						
	Were specified samp all analytical needs?	le aliquots obta	ined meeting	Are fewer than 3 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/COPECs ^c identified in another sample type?	If no single sample types being compared was significantly different in the number of COPCs/COPEcs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?				
	Attempt 1	Attempt 2	Attempt 3								
Whole Water	No	Yes	NA	Yes	15	Yes	NA				
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	16	No	No				
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	17	No	No				
LSM dissolved	No	Yes	NA	Yes	16	Yes	NA				
HSM dissolved	No	Yes	NA	Yes	14	Tes	NA				
LSM particulate	No	Yes	NA	Yes	13	Yes	NA				
HSM particulate	No	Yes	NA	Yes	16	103	NA				

Positive Target Analyte Identification and Concentration Comparison

Analyte Identified	PR1CSOCLYWW-02B Whole Water ^g (µg/L)	LQ ^h	VQ	PR1CSOCLYLD-02B LSM Dissolved* (µg/L)	LQ ^h	VQ	PR1CSOCLYHD-02B HSM Dissolved ^g (µg/L)	LQ ^h	VQ	% RPD	PR1CSOCLYLP-02B LSM Particulate ⁸ (μg/kg)	LQ ^h	VQ	PR1CSOCLYHP-02B HSM Particulate ⁶ (μg/kg)	LQ ^b	VQ	% RPD
Naphthalene				0.051	ВЈ		0.035	BD	J	37.2				90	BD	J	
2-Methylnaphthalene	0.044	D	J				0.052	D						76	D	М	
Acenaphthylene	0.0055	GD	J	0.0058	J		0.0025	GD		79.5	480	G	J				
Acenaphthene	0.013	D	J	0.014			0.015	D		6.90				52	D	M	
Fluorene	0.026	D	J	0.021			0.030	D		35.3				80	D	M	
Phenanthrene	0.065	D	J	0.038	В		0.064	D		51.0	2500	В	J	790	BD	M	104
Anthracene	0.013	D	J	0.015		No.	0.011	D		30.8	870		J	100	D	M	159
Fluoranthene	0.082	D	J	0.039	ВЈ		0.069	D		55.6	9100		J	1000	D	М	160
Pyrene	0.066	D	J	0.026	ВЈ	L	0.056	ם		73.2	8400		J	940	D	M	160
Benzo (a) anthracene	0.032	D	3	0.0074	Į	L	0.023	D		103	6700		J	580	Q	М	168
Chrysene	0.050	D	J	0.014	J	L	0.034	ם		83.3	8600		J	940	D	M	161
Benzo(b)fluoranthene	0.047	D	J	0.0081	J	L	0.033	D		121	7200		J	830	D	М	159
Benzo(k) fluoranthene	0.039	D	j	0.0061	J	L	0.029	ם		130	8500		J	750	D	M	168
Benzo(a) pyrene	0.030	D	J	0.0040	G J	L	0.020	D		133	6600		j	560	D	M	169
indeno(1,2,3-cd)pyrene	0.012	D	J	0.0021	G J	L					5100		J	540	D	М	162
Dibenzo (a, h) anthracene				0.00075	G J	L					1800		j	200	D	M	160
Benzo(g,h,i)perylene	0.012	D	j	0.0028	G J	L					6200		j	650	D	М	162
1-Methylnaphthalene	0.041	D	J	0.063	J		0.053	D		17.2				54	D	M	
Benzo[e]pyrene	0.031	D	J	0.0059	J	L	0.021	D		112	7300		j	650	D	М	167
Perylene	0.0089	D	J	0.00082	GJ	L	0.0054	GD		147	2000		j	170	D	М	169
3,6-Dimethylphenanthrene	0.0085	GD	J	0.0035	GB J	L	0.011	ם		103	500		J	53	D	M	162
1-Methylanthracene	0.016	D	J	0.0087			0.022	D		86.6	1700		J	110	ם	М	176
1-Methylfluoranthene	0.019	D	j	0.0072			0.016	D		75.9	2700		J	260	D	М	165
1-Methylpyrene	0.0063	GD	j	0.0024	G		0,0068	GD		95.7	840		J	74	D	М	168
2,6-Dimethylnaphthalene	0.069	D	J	0.053	J		0.092	D		53.8				70	D	M	
2,3,5-Trimethylnaphthalene	0.044	D	J	0.036	J		0.052	D	J	36.4	350	G	J	53	D	M	147
Dibenzofuran				0.0073			0.016	D		74.7				48	D	М	
1-Methylphenanthrene	0.025	D	j	0.0069			0.036	D		136	400	G	J	94	D	M	124
Dibenzothiophene	0.011	D	J	0.011			0.018	D		48.3				52	D	M	

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study
HSM = high-solids mass
LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions RPD = relative percent difference SIM = selective ion monitoring SVOC = semivolatile organic compound µg/L = micrograms per liter

µg/kg = micrograms per kilograms
VQ = laboratory qualifier - See Attachment 2 for definitions
% = percent

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

[°] COPCs/COPECs listed in the FFS: Naphthalene, Fluorene, Pyrene, Benzo(k)fluoranthene, Benzo(g,h,i)perylene, 2-methylnaphthalene, Phenanthrene, Benzo(a)anthracene,

Benzo(a)pyrene, Acenaphthylene, Anthracene, Chrysene, Indeno(1,2,3-cd)pyrene, Acenaphthene, Fluoranthene, Benzo(b)fluoranthene, and Dibenzo(a,h)anthracene.

d At least 2 more

e Fewer than 3

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^g No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 2 FIELD DUPLICATE - SEMIVOLATILES-SIM PR1**DUP-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

SVOC SIM Sample Collection Techniques	Sample C	ollection Quali	ty ^a	Analytical Quality ^b	ldentifica:	tion of Target Analy	tes
	Were specified samp meeting all analytica		ained	Are fewer than 3 results "R" qualified (rejected due to association with severe data quality issues)?	Number of COPCs/COPECs ^c listed in the FFS identified?	Are at least 2 more COPCs/COPECs ^c identified in another sample type?	If no single sample types being compared was significantly different in the number of COPCs/COPECs identified (distinguished by a single "no" in the previous column), are the overall number of target analytes identified significantly different?
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	No	Yes	NA	Yes	17	No	No
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	16	No	No
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	17	No	No
LSM dissolved	No	Yes	NA	Yes	15	Yes	NA
HSM dissolved	No	Yes	NA	Yes	13	ies	NA
LSM particulate	No	Yes	NA	Yes	14	Yes	NA
HSM particulate	No	Yes	NA	Yes	16	les .	NA

Positive Target Analyte Identification and Concentration Comparison f

Analyte Identified	PR1WWDUP-02B Whole Water ^g (µg/L)	LQ ^h	VQ	PR1LDDUP-02B LSM Dissolved ^g (µg/L)	LQ ^h	VQ	PR1HDDUP-02B HSM Dissolved ⁶ (µg/L)	LQ ^h	VQ	% RPD	PR1LPDUP-02B LSM Particulate ⁸ (µg/kg)	LQ ^h	VQ	PR1HPDUP-02B HSM Particulate ^s (µg/kg)	ια ^h	VQ	% RPD
Naphthalene	0.23	DB J		0.037	В	JL								410	BD	J	
2-Methylnaphthalene	0.25	D J	1				0.049	D						73	D	М	
Acenaphthylene	0.057	GD J	1	0.018		JL	0.003	GD		143	2500	GD	J				
Acenaphthene	0.12	D J	l	0.0072		JL.	0.013	D		57.4				40	D	M	
Fluorene	0.18	D J	1	0.014		JL	0.028	D		66.7	6900	D	J	66	D	М	196
Phenanthrene	1.5	D J	1	0.044	В	JL	0.060	D		30.8	65000	DB	J	590	BD	M	196
Anthracene	0.29	D J	1	0.012		JL	0.0089	D		29.7	10000	٥	J	82	D	M	197
Fluoranthene	2.9	D J	l	0,031	В	J	0.060	D		63.7	130000	۵	J	1100	O	М	197
Pyrene	1.8	D J		0.019	В		0.058	D		101	91000	۵	J	810	D	M	196
Benzo(a)anthracene	1.2	D J	1	0.0033	G		0.020	D		143	54000	D	J	470	D	М	197
Chrysene	1.7	D J	1	0.0083			0.032	D		118	83000	D	J	770	D	М	196
Benzo(b)fluoranthene	1.8	D J	1	0.0035	U		0.032	D		161	82000	D	J	720	D	М	197
Benzo(k)fluoranthene	1.3	D J	1	0.0022	G		0.026	D		169	64000	D	J	630	Q	М	196
Benzo(a)pyrene	1.3	D J	1	0.0018	Ø		0.018	D		164	56000	D	J	470	D	M	197
Indeno(1,2,3-cd)pyrene	1.1	D J	1	0.0010	G						44000	D	J	420	D	М	196
Dibenzo(a,h)anthracene	0.38	D J	1	477							16000	D	J	150	D	М	196
Benzo(g,h,i)perylene	1.3	D J	1	0,0016	G						55000	D	J	540	D	М	196
1-Methylnaphthalene	0.17	D J		0.034		JL	0.047	D		32.1				49	D	М	
Benzo[e]pyrene	1.3	D J	1	0.0026	U		0.019	D		152	61000	٥	J	570	D	М	196
Perylene	0.38	D J		0.00051	G		0.0058	GD		168	15000	٥	J	140	٥	М	196
3,6-Dimethylphenanthrene	0.13	D J		0.0028	GB	JL	0.0095	D		109	4300	GD	J	37	D	М	197
1-Methylanthracene	0.27	D J	l	0.0049		JL	0.016	D		106	15000	٥	J	80	O	М	198
1-Methylfluoranthene	0.46	D J	l	0.0036	G		0.013	D		113	24000	D	J	210	O	М	197
1-Methylpyrene	0.13	D J	1	0.0014	Ø		0.0061	GD		125	7100	D	J	64	D	М	196
2,6-Dimethylnaphthalene	0.21	D J	1	0.027		JL	0.087	D		105	5400	D	J	77	D	M	194
2,3,5-Trimethylnaphthalene	0.18	D J		0.014		JL	0.011	D	J	24.0	7500	GD	J	60	D	М	197
1,1'-Biphenyl				0.0049		JL											
Dibenzofuran	0.12	D J		0.0046		JL	0.0094	D		68.6				37	D	М	200
1-Methylphenanthrene	0.14	D J		0.0057		JL	0.032	D		140	10000	D	J	120	D	М	195
Dibenzothiophene	0.13	D J		0.015		JL	0.016	D		6.45	3700	GD	J	32	GD	М	197

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
FFS = focused fesability study
HSM = high-solidsmass
LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions
RPD = relative percent difference
SIM = selective ion monitoring
SVOC = semivolatile organic compound
µg/L = micrograms per liter

 $\mu g/kg$ = micrograms per kilograms VQ = laboratory qualifier - See Attachment 2 for definitions % = percent

 $^{^{\}rm b}{\rm Analytical}$ quality is based upon the program 90% analytical completeness objectives.

^c COPCs/COPECs listed in the FFS: Naphthalene, Fluorene, Pyrene, Benzo(k)fluoranthene, Benzo(g,h,i)perylene, 2-methylnaphthalene, Phenanthrene, Benzo(a)anthracene, Benzo(a)pyrene, Acenaphthylene, Anthracene, Chrysene, Indeno(1,2,3-cd)pyrene, Acenaphthene, Fluoranthene, Benzo(b)fluoranthene, and Dibenzo(a,h)anthracene.

^d At least 2 more

e Fewer than 3

f Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

g No rejected data.

h A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.



Detailed Evaluation Sheets (Worksheet #11) – Chlorinated Herbicides

EVENT 1 ATTEMPT 2 ORIGINAL SAMPLE - CHLORINATED HERBICIDES PR1SCOCLY**-01B

QAPP Worksheet #11-1

Project Quality Objectives/Systematic Planning Process Statements (Phase I)

Phase I Data Comparison Chart

Chlorinated Herbicides Sample Collection Technique	Sample Coll	ection Qualit	y ^a	Analytical Quality ^b		tion of Target nalytes
	Were specified sample meeting all analytical r		ined	Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	Yes	Yes	Yes	Yes	0	No
LSM dissolved plus LSM particulate	Yes	Yes	Yes	Yes	2	Yes
HSM dissolved plus HSM particulate	Yes	Yes	Yes	Yes	1	No
LSM dissolved	Yes	Yes	Yes	Yes	2	Yes
HSM dissolved	Yes	Yes	Yes	Yes	0	No
LSM particulate	Yes	Yes	Yes	Yes	0	No
HSM particulate	No	Yes	Yes	Yes	1	Yes

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1SCOCLYWW-01B Whole Water ^d (µg/L)	VQ	PR1CSOCLYLD-01B LSM Dissolved ^d (µg/L)	ເດ້	VQ	PR1CSOCLYHD-01B HSM Dissolved ^d (µg/L)	LQ* VQ	PR1CSOCLYLP-01B LSM Particulate (µg/kg)	PR1CSOCLYHP-01B HSM Particulate ^d (μg/kg)		VQ % RPD
2,4-DB			0.45		NJ						
2,4,5-T									24	G	JL
Silvex (2,4,5-TP)			0.02		J						

There are no COPCs/COPECs in the target analyte list for chlorinated herbicides.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern

HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

RPD = relative percent difference $\mu g/L$ = micrograms per liter $\mu g/kg$ = micrograms per kilograms

VQ = validation qualifier - See Attachment 2 for definitions

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

c This target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^d No rejected data.

e A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 1 ATTEMPT 2 FIELD DUPLICATE - CHLORINATED HERBICIDES PR1**DUP-01B

QAPP Worksheet #11-1

Project Quality Objectives/Systematic Planning Process Statements (Phase I)

Phase I Data Comparison Chart

Chlorinated Herbicides Sample Collection Technique	Sample Co	ollection Qua	lity ^a	Analytical Quality ^b	1	ation of Target nalytes
	Were specified sam meeting all analytic		btained	Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	Yes	Yes	Yes	Yes	0	No
LSM dissolved plus LSM particulate	Yes	Yes	Yes	Yes	1	Yes
HSM dissolved plus HSM particulate	Yes	Yes	Yes	Yes	1	Yes
LSM dissolved	Yes	Yes	Yes	Yes	1	Yes
HSM dissolved	Yes	Yes	Yes	Yes	0	No
LSM particulate	Yes	Yes	Yes	Yes	0	No
HSM particulate	No	Yes	Yes	Yes	1	Yes

Positive Target Analyte Identification and Concentration Comparison^c

	PR1WWDUP-01B			PR1LDDUP-01B						PR1LPDUP-01B	PR1HPDUP-01B	
	Whole Water ^d			LSM Dissolved ^d			PR1HDDUP-01B HSM			LSM Particulate ^d	HSM Particulate ^d	
Analyte Identified	(μg/L)	LQ ^e	VQ	(μg/L)	LQ*	va	Dissolved ^d (µg/L)	rd, Ad	% RPD	(μg/kg) LQ ^e	VQ (μg/kg)	LQ" VQ % RPD
2,4-DB				1		NJ						
2,4,5-T											140	G J

There are no COPCs/COPECs in the target analyte list for chlorinated herbicides.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern

HSM = high-solidsmass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

RPD = relative percent difference μg/L = micrograms per liter μg/kg = micrograms per kilograms

VQ = validation qualifier - See Attachment 2 for definitions

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

 $^{^{\}rm b}$ Analytical quality is based upon the program 90% analytical completeness objectives.

^c This target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

d No rejected data.

and a "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

EVENT 2 ORIGINAL SAMPLE - CHLORINATED HERBICIDES PR1CSOCLY**-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Chlorinated Herbicides Sample Collection Technique	Sample Coli	ection Qualit	y [°]	Analytical Quality ^b		tion of Target nalytes
	Were specified sample meeting all analytical n		ined	Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	No	Yes	NA	Yes	0	No
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	0	No
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	1	Yes
LSM dissolved	No	Yes	NA	Yes	0	No
HSM dissolved	No	Yes	NA	Yes	1	Yes
LSM particulate	No	Yes	NA	Yes	0	No
HSM particulate	No	Yes	NA	Yes	0	No

Positive Target Analyte Identification and Concentration Comparison

	PR1CSOCLYWW-02B Whole Water ^d (μg/L)	LQ°	VQ	PR1CSOCLYLD-02B LSM Dissolved d (µg/L)	LQ*	8	PR1CSOLCYHD-02B HSM Dissolved ^d (μg/L)	ια° vα	% RPD	PR1CSOCLYLP-02B LSM Particulate ^d (µg/kg)	۷Q	PR1CSOCLYHP-02B HSM Particulate ^d (μg/kg)	% RPD
2,4-DB				Constant of the Constant of th			0.31	B NJ					

There are no COPCs/COPECs in the target analyte list for chlorinated herbicides.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern

HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

RPD = relative percent difference

μg/L = micrograms per liter

μg/kg = micrograms per kilograms

VQ = validation qualifier - See Attachment 2 for definitions

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt,

 $^{^{\}mathrm{b}}$ Analytical quality is based upon the program 90% analytical completeness objectives.

^c This target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^d No rejected data.

e A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitationely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 2 FIELD DUPLICATE - CHLORINATED HERBICIDES PR1**DUP-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Chlorinated Herbicides Sample Collection Technique	Sample Co	ollection Qua	lity ^a	Analytical Quality ^b		ation of Target nalytes
	Were specified sam meeting all analytic		bbtained	, ,	Number of target analytes identified?	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	No	Yes	NA	Yes	0	No
LSM dissolved plus LSM particulate	No	Yes	NA	Yes	2	Yes
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	0	No
LSM dissolved	No	Yes	NA	Yes	2	Yes
HSM dissolved	No	Yes	NA	Yes	0	No
LSM particulate	No	Yes	NA	Yes	0	No
HSM particulate	No	Yes	NA	Yes	0	No

Positive Target Analyte Identification and Concentration Comparison

Analyte Identified	PR1WWDUP-02B Whole Water ^d (µg/L)	ĽQ [€]	VQ	PR1LDDUP-02B LSM Dissolved ^d (µg/L)	VQ	PR1HDDUP-02B HSM Dissolved ^d (µg/L) LQ ^e VQ	88	PR1LPDUP-02B LSM Particulate ^d (µg/kg) LQ ^e	PR1HPDUP-02B HSM Particulate ^d (µg/kg) LQ ^e VQ % RPD
2,4-DB				0.41	NJ				
2,4,5-T		·	·	0.21					

There are no COPCs/COPECs in the target analyte list for chlorinated herbicides.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LSM = low-solids mass
LQ = laboratory qualifier - See Attachment 1 for definitions

RPD = relative percent difference μg/L = micrograms per liter μg/kg = micrograms per kilograms VQ = validation qualifier - See Attachment 2 for definitions

a Na" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c This target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

d No rejected data.

e A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

EVENT 1 ATTEMPT 3 ORIGINAL SAMPLE - CHLORINATED HERBICIDES PR1CSOCLY**-01C

QAPP Worksheet #11-1

Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

Chlorinated Herbicides Sample Collection Technique	Sample Colle	ection Quality		Analytical Quality ^b		ation of Target nalytes
	Were specified sample ali analytical needs?	quots obtaine	d meeting all	Is fewer than 1 result "R" qualified (proceed due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	Yes	Yes	Yes	Yes	4	Yes
LSM dissolved plus LSM particulate	Yes	Yes	Yes	Yes	2	No
HSM dissolved plus HSM particulate	Yes	Yes	Yes	Yes	4	Yes
LSM dissolved	Yes	Yes	Yes	Yes	2	No
HSM dissolved	Yes	Yes	Yes	Yes	4	Yes
LSM particulate	Yes	Yes	Yes	Yes	0	No
HSM particulate	No	Yes	Yes	Yes	0	No

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1CSOCLYWW-01C Whole Water ^d (µg/L)	LQ ^e	VQ	PR1CSOCLYLD-01C LSM Dissolved ^d (µg/L)	ſŎ _ŧ	VQ	PR1CSOCLYHD-01C HSM Dissolved ^d (µg/L)	LQ°	VQ	% RPD	PR1CSOCLYLP-01C LSM Particulate ^d (μg/kg)	LQ°	VQ	PR1CSOCLPHP-01C HSM Particulate ^d (μg/kg)	/Q % RPD
2,4-D	0.36	В	NJ	0.47	В		0.40	В		16.1					
2,4-DB	0.59	В		7			0.47	В	ΝJ						
2,4,5-T	0.10	G	NJ	0.09	G	NJ	0.022	G	NJ	123					
Silvex (2,4,5-TP)	0.051	В					0.023	В							

There are no COPCs/COPECs in the target analyte list for chlorinated herbicides.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solidsmass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

RPD = relative percent difference µg/L = micrograms per liter µg/kg = micrograms per kilograms

VQ = validation qualifier - See Attachment 2 for definitions

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

c This target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

^d No rejected data.

e A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

EVENT 1 ATTEMPT 3 DUPLICATE SAMPLE - CHLORINATED HERBICIDES PR1**DUP-01C

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Chlorinated Herbicides Sample Collection Technique	Sample Co	llection Quali	ty³	Analytical Quality ^b		ation of Target nalytes
	Were specified samp meeting all analytical		tained	Is fewer than 1 result "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes identified?	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	Yes	Yes	Yes	Yes	4	Yes
LSM dissolved plus LSM particulate	Yes	Yes	Yes	Yes	4	Yes
HSM dissolved plus HSM particulate	Yes	Yes	Yes	Yes	3	No
LSM dissolved	Yes	Yes	Yes	Yes	4	Yes
HSM dissolved	Yes	Yes	Yes	Yes	3	No
LSM particulate	Yes	Yes	Yes	Yes	0	No
HSM particulate	No	Yes	Yes	Yes	0	No

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1WWDUP-01C Whole Water ^d (µg/L)	LQ [€]		PR1LDDUP-01C LSM Dissolved ^d (μg/L)	ιq°	VQ	PR1HDDUP-01C HSM Dissolved ^d (µg/L)		VQ	% RPD	PR1LPDUP-01C LSM Particulate ^d (µg/kg)	LQ*	PR1HPDUP-01C HSM Particulate ^d (µg/kg)	vq 9	% RPD
2,4-D	0.48	В		0.51	В	JH	0.41	В		21.7					
2,4-DB	0.28	В	NJ	0.44	В	NJ									
2,4,5-T	0.1		NJ	0.07	G	NJ	0.054	G	NJ	31.3					
Silvex (2,4,5-TP)	0.032	В	NJ	0.021	В	JH	0.021	В	lИ	0.00					

There are no COPCs/COPECs in the target analyte list for chlorinated herbicides.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass

LSM = low-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

RPD = relative percent difference µg/L = micrograms per liter µg/kg = micrograms per kilograms

VQ = validation qualifier - See Attachment 2 for definitions

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

 $^{^{\}rm b}$ Analytical quality is based upon the program 90% analytical completeness objectives.

c This target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list. Additional pages may be necessary.

d No data rejected

e A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

	pp				R.A
4	nn	Δn	m	ľ	IVI
, ,	\sim			_	

Detailed Evaluation Sheets (Worksheet #11) - Cyanide

EVENT 1 ORIGINAL SAMPLE - CYANIDE PR1CSOCLY**-01B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

Cyanide Sample Collecton Techniques	Sample Co	ollection Quality	ı	Analytical Quality ^b	Identification of Target Analytes		
II .	Were specified sample aliquots obtained meeting all f				Was cyanide positively identified?		
	Attempt 1 Attempt 2 Attempt 3						
Whole Water	Yes	Yes	NA	Yes	Yes (1)		
HSM dissolved plus HSM particulate ^c	No	Yes	NA	Yes	Yes (1)		

Positive Target Analyte Identification and Concentration Comparison^d

Analyte Identified	PR1CSOCLYWW-01B Whole Water ^e Concentration (µg/L)	VQ	PR1CSOCLYHD-01B HSM Dissolved ^e Concentration (µg/L)	LQ	VQ	PR1CSOCLYHP-01B HSM Particulate ^{ce} Concentration (mg/Kg)	LQ	VQ
Cyanide	29.3		31.3			5.8		J

There are no COPC/COPECs in the target list for Cyanide.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern HSM = high-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

μg = micrograms

VQ = laboratory qualifier - See Attachment 2 for definitions

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c HSM particulate based on a composite of debris and fines.

^d Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^e No rejected data.

EVENT 1 FIELD DUPLICATE - CYANIDE PR1**DUP-01B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

Cyanide Sample Collecton Techniques	Sample Collect	tion Quality ^a		Analytical Quality ^b	Identification of Target Analytes
	Were specified sample aliquots obtained meeting all			Is the cyanide result free of any "R" flag (rejected due to association with severe data quality issues)?	Was cyanide positively identified?
	Attempt 1	Attempt 2	Attempt 3		
Whole Water	Yes Yes NA		Yes	Yes (1)	
HSM dissolved plus HSM particulate ^c	No	Yes	NA	Yes	Yes (1)

Positive Target Analyte Identification and Concentration Comparison^d

	Analyte Identified	PR1WWDUP-01B Whole Water ^e Concentration (μg/L)	LQ	VQ	PR1HDDUP-01B HSM Dissolved ^e Concentration (µg/L)	LQ	VQ	PR1HPDUP-01BHSM Particulate ^{ce} Concentration (mg/Kg)	LQ	VQ
С	'yanide	27.2			31.6			6.4		J

There are no COPC/COPECs in the target list for Cyanide.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

HSM = high-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

μg = micrograms

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c HSM particulate based on a composite of debris and fines.

^d Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^e No rejected data.

EVENT 2 ORIGINAL SAMPLE - CYANIDE PR1CSOCLY**-02B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

Cyanide Sample Collecton Techniques	Sample Coll	ection Quality	y ^a	Analytical Quality ^b	Identification of Target Analytes
	Were specified sample aliquots obtained fla		Is the cyanide result free of any "R" flag (rejected due to association with severe data quality issues)?	Was cyanide positively identified?	
	Attempt 1	Attempt 2	Attempt 3		
Whole Water	No Yes NA		Yes	Yes (1)	
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	Yes (1)

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1CSOCLYWW-02B Whole Water ^d Concentration (μg/L)	LQ	VQ	PR1CSOCLYHD-02B HSM Dissolved ^d Concentration (µg/L)	LQ VC	PR1CSOCLYHP-02B HSM Particulate ^d Concentration (mg/Kg)	VQ
Cyanide	3.8	В	j	ND	U	2.4	M

There are no COPC/COPECs in the target list for Cyanide.

Notes:

COPCs = contaminants of potential concern COPECs = contaminants of potential ecological concern HSM = high-solids mass LQ = laboratory qualifier - See Attachment 1 for definitions

μg = micrograms

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^d No rejected data.

EVENT 2 FIELD DUPLICATE - CYANIDE PR1**DUP-02B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

Cyanide Sample Collecton Techniques	Sample Col	lection Quali	ty ^a	Analytical Quality ^b	Identification of Target Analytes
	"R" Were specified sample aliquots obtained ass		Is the cyanide result free of any "R" flag (rejected due to association with severe data quality issues)?	Was cyanide positively identified?	
	Attempt 1	Attempt 2	Attempt 3		
Whole Water	No Yes NA		Yes	Yes (1)	
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	Yes (1)

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1WWDUP-02B Whole Water ^d Concentration (µg/L)	LQ	VQ	PR1HDDUP-02B HSM Dissolved ^d Concentration (μg/L)	LQ	να	PR1HPDUP-02B HSM Particulate ^d Concentration (mg/Kg)	LQ	VQ
Cyanide	2.3	В	j	ND		U	1.6		

There are no COPC/COPECs in the target list for Cyanide.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

μg = micrograms

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^d No rejected data.

Appendix N

Detailed Evaluation Sheets (Worksheet #11) - VOCs

EVENT 1 ORIGINAL SAMPLE - VOLATILE ORGANIC COMPOUND PR1CSOCLY**-01B

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

VOC Sample Collecton Techniques	Sample Collection Quality ^a			Analytical Quality ^b	Identification of Target Analytes				
	Were specified sample aliquots obtained meeting (Are fewer than 2 results "R" qualified (rejected due to association with severe data quality issues)?		Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?			
	Attempt 1	Attempt 2	Attempt 3						
Whole Water	Yes	Yes	NA	Yes	1	Yes			
HSM dissolved plus HSM particulate $^{\circ}$	No	Yes	NA	No (4) ^d	NA	NA			

Positive Target Analyte Identification and Concentration Comparison^e

Analyte Identified	PR1CSOCLYWW-01B Whole Water ^f (μg/L)	LQ ^g	VQ	PR1CSOCLYHD-01B HSM Dissolved ^f (µg/L)	LQ ^g	VQ	PR1CSOCLYHP-01B HSM Particulate ^c (μg/Kg)	LQ ^g	VQ
1,4-Dichlorobenzene	0.24	G		0.21	G		47		J
Chlorobenzene				100 100 100 100 100 100 100 100 100 100			1.4	G	J
1,,3-Dichlorobenzene							ND	U	R ^h
1,2-Dichlorobenzene							ND	U	R ^h
1,2,4-Trichlorobenzene							ND	U	R^h
1,2,3-Trichlorobenzene							ND	U	R ^h

There are no COPC/COPECs in the target list for VOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LQ = laboratory qualifier - See Attachment 1 for definitions
R = rejected data result

µg/L = micrograms per liter µg/Kg = micrograms per kilogram VQ = laboratory qualifier - See Attachment 2 for definitions VQ = validation qualifier

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^bAnalytical quality is based upon the program 90% analytical completeness objectives.

^c HSM particulate based on a composite of debris and fines.

 $^{^{\}rm d}$ Values in paretheses indicate the total number of rejected results.

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^f No rejected data.

⁸ A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitationally less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

h PRICSOCLYHP-01B Data results rejected due to low internal standard recovery. Sample not used during sample collection technique evaluation.

EVENT 1 FIELD DUPLICATE - VOLATILE ORGANIC COMPOUND PR1**DUP-01B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

VOC Sample Collecton Techniques	Sample Co	llection Quali	i+v ^a	Analytical Quality ^b	Identification of T	-argot Δnalvtos
	Sample Collection Quality ^a			Analytical Quanty	I Adentification of a	I argue Ariunytes
	q Were specified sample aliquots obtained a		Are fewer than 2 results "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?	
	Attempt 1 Attempt 2 Attempt 3					
Whole Water	Yes Yes NA		Yes	1	Yes	
HSM dissolved plus HSM particulate ^c	No	Yes	NA	No (4) ^d	NA	NA

Positive Target Analyte Identification and Concentration Comparison^e

Analyte Identified	PR1WWDUP-01B Whole Water ^f (μg/L)	LQ ^g	VQ	PR1HDDUP-01B HSM Dissolved ^f (µg/L)	LQ [#]	VQ	PR1HPDUP-01B HSM Particulate ^c (µg/Kg)	LQ ^g	VQ
1,4-Dichlorobenzene	0.22	G		0.22	G		15		J
Chlorobenzene				1000			0.5	G	J
1,3-Dichlorobenzene							ND	U	R ^h
1,2-Dichlorobenzene							ND	U	R ^h
1,2,4-Trichlorobenzene							ND	U	R ^h
1,2,3-Trichlorobenzene							ND	U	R ^h

There are no COPC/COPECs in the target list for VOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LQ = laboratory qualifier - See Attachment 1 for definitions
R= rejected data result

 μ g/L = micrograms per liter μ g/Kg = micrograms per kilogram VQ = laboratory qualifier - See Attachment 2 for definitions VQ = validation qualifier

a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c HSM particulate based on a composite of debris and fines.

 $^{^{\}rm d}$ Values in paretheses indicate the total number of rejected results.

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^f No rejected data

^g A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

h PR1HPDUP-01B Data results rejected due to low internal standard recovery. Sample not used during sample collection technique evaluation.

EVENT 2 ORIGINAL SAMPLE - VOLATILE ORGANIC COMPOUND PR1CSOCLY**-02A

QAPP Worksheet #11-1
Project Quality Objectives/Systematic Planning Process Statements (Phase I)
Phase I Data Comparison Chart

VOC Sample Collecton Techniques	Sample Colle	ction Quality	à	Analytical Quality ^b	Identification of Target Analytes				
II .	Were specified sample aliquots obtained meeting					Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?			
	Attempt 1	Attempt 2	Attempt 3						
Whole Water	Yes NA NA		Yes	1	Yes				
HSM dissolved plus HSM particulate ^c	Yes NA NA			No (5) ^d	NA	NA			

Positive Target Analyte Identification and Concentration Comparison^e

Analyte Identified	PR1CSOCLYWW-02A Whole Water ^f (µg/L)	LQ ^g	vq	PR1CSOCLYHD-02A HSM Dissolved ^f (µg/L)	LQ ^g	۷q	PR1CSOCLYHP-02A2 HSM Particulate ^c (μg/Kg)	LQ ^g	VQ
1,4-Dichlorobenzene	0.079	G		0.081	G				
Chlorobenzene							ND	U	R ^h
1,3-Dichlorobenzene							ND	٦	R ^h
1,4-Dichlorobenzene							ND	U	R ^h
1,2-Dichlorobenzene							ND	U	R^h
1,2,3-Trichlorobenzene				Total Comments (Indiana)			ND	U	R^h

There are no COPC/COPECs in the target list for VOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LQ = laboratory qualifier - See Attachment 1 for definitions
R = rejected data result

μg/L = micrograms per liter μg/Kg = micrograms per kilogram VQ = laboratory qualifier - See Attachment 2 for definitions VQ = validation qualifier

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^cHSM particulate based on a composite of debris and fines.

^d Values in paretheses indicate the total number of rejected results.

e Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^f No rejected data

⁸ A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitationally less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

h PR1CSOCLYHP-02A2 Data results rejected due to low internal standard recovery. Sample not used during sample collection technique evaluation.

EVENT 2 FIELD DUPLICATE - VOLATILE ORGANIC COMPOUND PR1** DUP-02A

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

VOC Sample Collecton Techniques	Sample Colle	ection Quality	a	Analytical Quality ^b	Identification of 1	Farget Analytes
	Were specified sample a all analytical needs?	liquots obtair		Are fewer than 2 results "R" qualified (rejected due to association with severe data quality issues)?	Number of target analytes	Does the sample collection technique have at least one more target analyte identified than the other sample collection technique?
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	Yes	NA	NA	Yes	1	Yes
HSM dissolved plus HSM particulate	Yes	NA	NA	No (4) ^c	NA	NA

Positive Target Analyte Identification and Concentration Comparison ^d

Analyte Identified	PR1WWDUP-02A Whole Water ^e (µg/L)	LQ ^f	VQ	PR1CSOCLYHD-02A HSM Dissolved ^e (µg/L)	ιQ ^f	VQ	PR1HPDUP-02A2 HSM Particulate (μg/Kg)	ια ^f	vq
1,4-Dichlorobenzene	0.080	G		0.078	G				
1,3-Dichlorobenzene							ND	U	R ^g
1,4-Dichlorobenzene							DN	U	R ⁸
1,2-Dichlorobenzene							ND	U	R ^g
1,2,3-Trichlorobenzene		·					ND	U	R ^g

There are no COPC/COPECs in the target list for VOCs.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LQ = laboratory qualifier - See Attachment 1 for definitions
R = rejected data result

 $\mu g/L =$ micrograms per liter $\mu g/Kg =$ micrograms per kilogram VQ = laboratory qualifier - See Attachment 2 for definitions VQ = validation qualifier

a NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c Values in paretheses indicate the total number of rejected results.

d Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^e No rejected data

f A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate. Values associated with a "G" qualifier are quantitatiovely less certain than those not associated with a "G" qualifier. This is because "G" qualified results fall below the low point of the calibration curve.

⁸ PR1HPDUP-02A2 Data results rejected due to low internal standard recovery. Sample not used during sample collection technique evaluation.

Appendix O

Detailed Evaluation Sheets (Worksheet #11) – TEPH

EVENT 1 ORIGINAL - TOTAL EXTRACTABLE PETROLEUM HYDROCARBONS PR1CSOCLY**-01B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

TEPH Sample Collecton Techniques	Sample Collection Quality ^a			Analytical Quality ^b	Identification of Target Analytes		
	Were specified sample ali all analytical needs?	quots obtaine		Is the TEPH result free of any "R" flag (rejected due to association with severe data quality issues)?	Was TEPH positively identified?		
	Attempt 1	Attempt 2	Attempt 3				
Whole Water	Yes	Yes	NA	Yes	Yes (1)		
HSM dissolved plus HSM particulate ^c	No	Yes	NA	Yes	Yes (1)		

Positive Target Analyte Identification and Concentration Comparison^d

Analyte Identified	PR1CSOCLYWW-01B Whole Water ^e Concentration (mg/L)	LQ	VQ	PR1CSOCLYHD-01B HSM Dissolved ^e Concentration (mg/L)	LQ	VQ	PR1CSOCLYHP-01B HSM Particulate ^{c,e} Concentration (mg/Kg)	LQ	να
ТЕРН	5.0	В	j	5.6	В	j	13000	BD	J

There are no COPC/COPECs in the target list for TEPH.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LQ = laboratory qualifier - See Attachment 1 for definitions

mg = milligrams
TEPH = total extractable petroleum hydrocarbon

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c HSM particulate based on a composite of debris and fines.

^d Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

e No rejected data

EVENT 1 DUPLICATE- TOTAL EXTRACTABLE PETROLEUM HYDROCARBONS PR1**DUP-01B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

TEPH Sample Collecton Techniques	Sample Collection Quality ^a			Analytical Quality ^b	Identification of Target Analytes
	Were specified sample aliquots obtained			Is the TEPH result free of any "R" flag (rejected due to association with severe data quality issues)?	Was TEPH positively identified?
	Attempt 1	Attempt 2	Attempt 3		
Whole Water	Yes	Yes	NA	Yes	Yes (1)
HSM dissolved plus HSM particulate ^c	No	Yes	NA	Yes	Yes (1)

Positive Target Analyte Identification and Concentration Comparison^d

Analyte Identified	PR1WWDUP-01B Whole Water ^e Concentration (mg/L)	LQ	VQ	PR1HDDUP-01B HSM Dissolved e Concentration (mg/L)	LQ	VQ	PR1HPDUP-01B HSM Particulate ^{c,e} Concentration (mg/Kg)	ıQ	VQ
ТЕРН	7.7	BD	***	3.5	В	J	13000	BD	J

There are no COPC/COPECs in the target list for TEPH.

Notes:

COPCs = contaminants of potential concern
COPECs = contaminants of potential ecological concern
HSM = high-solids mass
LQ = laboratory qualifier - See Attachment 1 for definitions

mg = milligrams

TEPH = total extractable petroleum hydrocarbon

VQ = laboratory qualifier - See Attachment 2 for definitions

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c HSM particulate based on a composite of debris and fines.

^d Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^e No rejected data

EVENT 2 ORIGINAL - TOTAL EXTRACTABLE PETROLEUM HYDROCARBONS PR1CSOCLY**-02B

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I)

Phase I Data Comparison Chart

TEPH Sample Collecton Techniques	Sample Collection Quality ^a			Analytical Quality ^b	Identification of Target Analytes	
	Were specified sample al all analytical needs?	iquots obtain		Is the TEPH result free of any "R" flag (rejected due to association with severe data quality issues)?	Was TEPH positively identified?	
	Attempt 1	Attempt 2	Attempt 3			
Whole Water	No	Yes	NA	Yes	Yes (1)	
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	Yes (1)	

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PR1CSOCLYWW-02B Whole Water ^d Concentration (mg/L)	LQ	VQ	PR1CSOCLYHD-02B HSM Dissolved d Concentration (mg/L)	LQ	VQ	PR1CSOCLYHP-02B HSM Particulate ^d Concentration (mg/Kg)	LQ	VQ
ТЕРН	2.22	D	J	ND		U,J	13000	D	J

There are no COPC/COPECs in the target list for TEPH.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

HSM = high-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

mg = milligrams

TEPH = total extractable petroleum hydrocarbon

VQ = laboratory qualifier - See Attachment 2 for definitions

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^d No rejected data

EVENT 2 DUPLICATE - TOTAL EXTRACTABLE PETROLEUM HYDROCARBONS PRDUP-02B**

QAPP Worksheet #11-1 Project Quality Objectives/Systematic Planning Process Statements (Phase I) Phase I Data Comparison Chart

TEPH Sample Collecton Techniques	Sample Collection Quality ^a			Analytical Quality ^b	Identification of Target Analytes
	Were specified sample aliquots obtained fla		Is the TEPH result free of any "R" flag (rejected due to association with severe data quality issues)?	Was TEPH positively identified?	
	Attempt 1	Attempt 2	Attempt 3		
Whole Water	No	Yes	NA	Yes	Yes (1)
HSM dissolved plus HSM particulate	No	Yes	NA	Yes	Yes (1)

Positive Target Analyte Identification and Concentration Comparison^c

Analyte Identified	PRWWDUP-02B Whole Water ^d Concentration (mg/L)	LQ	VQ	PR1HDDUP-02B HSM Dissolved ^d Concentration (mg/L)	LQ	VQ	PR1HPDUP-02B HSM Particulate ^d Concentration (mg/Kg)	LQ	vq
ТЕРН	4.200		j	ND		U,J	7700	D	J

There are no COPC/COPECs in the target list for TEPH.

Notes:

COPCs = contaminants of potential concern

COPECs = contaminants of potential ecological concern

HSM = high-solids mass

LQ = laboratory qualifier - See Attachment 1 for definitions

mg = milligrams

TEPH = total extractable petroleum hydrocarbon

VQ = laboratory qualifier - See Attachment 2 for definitions

^a A "NA" in one of the Attempt columns indicates that the analytical group had already been collected in a previous attempt and was not intended to be collected during that column's attempt.

^b Analytical quality is based upon the program 90% analytical completeness objectives.

^c Positive target analyte identification and concentration comparison chart will comprise the detected analytes from the full target analyte list.

^d No rejected data

Appendix P CSO/SWO Phase I Data Quality Usability Assessment Report

Combined Sewer Overflow/Stormwater Outfall Investigation

Lower Passaic River Study Are

Phase I Data Quality Usability Assessment Report

Prepared For

Tierra Solutions, Inc.

East Brunswick, NJ

Prepared By

Environmental Data Services, Ltd.

June 2016

Revision 2

TABLE OF CONTENTS

I	Background	1
2	Introduction	1
3	Data Quality Parameters Overview	1
	3.1 Precision	2
	3.2 Accuracy/Bias Contamination	4
	3.3 Overall Accuracy/Bias	5
	3.4 Sensitivity	6
	3.5 Representativeness	15
	3.6 Comparability	16
	3.7 Completeness	17
4	Phase I CSO/SWO Investigation Data Verification/Validation	21
	4.1 Data Quality Issues	21
	4.1.1 Whole Water Samples Data Quality Issues	22
	4.1.2 Whole Water Samples Systematic and Random Data Quality Issues	
	by Analytical Group	22
	4.1.3 Low Solids Mass Samples Systematic Data Quality Issues	30
	4.1.4 Low Solids Mass Samples Systematic and Random Data Quality Issues	
	by Analytical Group	31
	4.1.5 High Solids Mass Samples Systematic Data Quality Issues	41
	4.1.6 High Solids Mass Samples Systematic and Random Data Quality Issues	
	by Analytical Group	41
	4.1.7 Grab Water Samples Systematic Data Quality Issues	57
	4.1.8 Grab Water Samples Systematic and Random Data Quality Issues	
	by Analytical Group	57
5	Total Tetrachlorinated Debenzo-p-Dioxin Verification	60
6	Conclusions	61
7	References	61

1. Background

In 2013 the United States Environmental Protection Agency (USEP A) approved a Quality Assurance Project Plan (QAPP) prepared by Tierra Solutions, Inc. (Tierra) for the investigation and characterization of combined sewer overflows (CSO s) and storm water outfalls (SW Os). The CSO/SWO Investigation QAPP, Revision 3 (Tierra, 2013) (hereafter referred to as the Q APP) outlined a two phased program – Phase I being a limited sampling effort with the objective of e valuating alternative sampling approaches and Phase II being a more fulsome sampling effort incorporating more overflows and outfalls.

The Phase I activities, conducted between June 10 th, 2013 and May 5 th, 2014, consisted of the collection and analysis of two CSO effluent samples using three approaches to sample collection: low solids mass (LSM), high solids mass (HSM) and whole water. Data collected will be evaluated to inform the selection of the most appropriate sampling approach to quantify contamina nts in the solid (particulate), dissolved, and whole water-phases during Phase II. The Phase I CSO effluent samples were collected at the Clay Street CSO location (described in Table 3-1 of the QAPP) and distributed to multiple laboratories for analyses. Validation of the sample analytical results was completed on July 14 th, 2014. According to Worksheet #33 of the QAPP, (Tierra, 2013) a Data Quality Usability Assessment Report (DQUAR) must be completed within 40 days of the conclusion of validation tasks.

2. Introduction

In accordance with requirements of the QAPP, the data quality usability assessment was conducted on both verified and validated data; this DQUAR provides a summary of the documentation and evaluation of data quality and usability for sample data collected during the implementation of Phase I of the CSO/SWO Investigation. The data verification and data validation processes are described respectively in Worksheets #34 and #35 of the QAPP. The information presented in this document will be used as part of the final Phase I evaluation that will determine the sampling method for each analytical group that will provide the greatest percentage of useable data to meet program data use and data quality objectives.

Worksheet #37 of the QAPP provides description of the components of the DQUAR. These components are described in detail in subsequent sections of this report.

3. Data Quality Parameters Overview

To assess whether the analytical data obtained were consistentwith the objectives of the QAPP, seven data quality parameters were evaluated. In the event that the data verification/validation process identified an instance where any of the data quality parameters did not meet the objectives established in the QAPP, the affected sample results were evaluated in accordance with the data verification/validation protocols specified in Worksheet #35 of the QAPP and documented accordingly. A detailed narrative describing the verification/validation assessments and findings can be found within the data verification/validation data assessment narratives prepared for each data package.

ine	seven data quality parameters assessed included the following:
	precision;
	accuracy/bias contamination;
	overall accuracy/bias;
	sensitivity;
	representativeness;
	comparability; and
	completeness.

Each of these data quality parameters, as it relates to Phase I of the QAPP program, is discussed below.

3.1 Precision

Precision is the measure of variability between individual sampe measurements of the same property under similar conditions. During the CSO/SWO Investigation program, precision was evaluated through the analysis of two types of duplicate samples. Field and laborato ry duplicates were analyzed at regular, specified intervals throughout the CSO/SWO Investigation program.

Field duplicates consisted of samples that were collected in the field at the frequency specified in the QAPP in order to determine the precision of field sampling methods. These samples were homogenized (except for those to be analyzed for volatile organic compounds [VOCs]) , split into two distinct samples, and submitted "blind" to the analytical laboratories for analysis (.e., the sample identification did not reveal the sample with which its field duplicate was associated).

Relative percent differences (RPDs) between the field sample results and the field duplicate results provide an estimate of the overall sampling and analytical precision.

Laboratory duplicates are two portions of a single homogeneous sample that are analyzed for the same parameter in order to determine the precision of the analytical system. Two types of laboratory duplicates were prepared. Laboratory duplicates without known analyte spi kes added were analyzed to monitor laboratory precision for cyanide, total organic carbon (TOC), t otal suspended solids (TSS), and total dissolved solids (TDS) analyses, while matrix spike (MS) and matrix spike duplicate (MSD) evaluations were performed to monitor laboratory precision for the remainin g analysis types. Laboratory duplicates were analyzed at the frequency specified in QAPP. The RPD betw een results obtained for a given laboratory duplicate pair provides an estimate of analytical precision.

The precision assessment for field and laboratory duplicate analyses is expressed as the RPD:

where: S = original sample concentration
D = duplicate sample concentration

Acceptance criteria for field and laboratory duplicates are pro vided in Worksheet #12 of the QAPP. Conformance to laboratory duplicate frequency requirements, aswell as acceptability of the resulting RPD values, were evaluated and considered during data validation.

Although laboratory duplicate analyses are used as indicators 6 relative precision of the analytical systems, the degree of homogeneity of the contaminants in the sample medum can also affect the reproducibility of a particular measurement. For example, pieces of decayed wood debris, chunks of asphalt, glass, free product, etc., can increase sample heterogeneity and thereforecan reduce the laboratory technician's ability to create homogeneous duplicate samples with which to measure p recision. Since the sample matrix characteristics can affect the way precision is measured, the s ample matrix should be considered by the validator.

With respect to the results of the Phase I CSO/SWO Investigation data, there are no limitations on data usage based on precision quality acceptance criteria. The following table summarizes the Phase I precision quality evaluation by analytical group and sampling technique. The "x" designation indicates that an issue was identified however, such issue does not infer that the data is unusable. A more detailed discussion of this data quality parameter evaluation is provided in Section 4.1 of this report.

Precision					
Analytical Groups	Whole Water	LSM	HSM	Grab Water	
Semivolatile Organics	X	X	X	-	
Volatile Organics (trace)		•	X	-	
Aroclor PCBs				-	
Organochlorine Pesticides		X	X	-	
Semivolatile Organics (SIM)		X	X	-	
Metals		-	-	X	
Mercury	X	-	-		
Methylmercury		-	-		
Cyanide		-	Х	-	
PCDD/PCDFs	X	X	X	-	
PCB Congeners	Х	X	Х	-	
Chlorinated Herbicides	Х	X	Х	-	
TOC/POC/DOC				-	
ТЕРН	Х	-	Х	-	
TSS	Х		Х		
TDS					
Grain Size		-	-	-	

^{- =} analysis was not performed for this analytical group

x = data qualified during validation for this analytical group

3.2 Accuracy/Bias Contamination

Accuracy parameters were also assessed with respect to contamin ation through the use of field and laboratory blanks. Any contamin ation present in field or labor atory blanks reflects the potential for contamination in associated samples. Measurement performance criteria for accuracy/bias contamination are outlined in Worksheet #12 of the QAPP. Acceptability of qulity control (QC) results for accuracy/bias contamination and conformance to field and laboratory QC samplefrequency requirements were evaluated and considered during the data verification/validation.

With respect to the results of the Phase I CSO/SWO Investigation data, there are no limitations on the data usage based on accuracy/bias contamination acceptance criteria. The following table summarizes the Phase I accuracy/bias contamination quality evaluation by analytical group and sampling technique. The "x" designation indicates that an issue was identified however, suc h issue does not infer that the data is unusable. A more detailed discussion of this data quality parameter evaluation is provided in Section 4.1 of this report.

Accuracy/Bias Contamination					
Analytical Groups	Whole Water	LSM	HSM	Grab Water	
Semivolatile Organics	X	X	X	-	
Volatile Organics (trace)		-	X	-	
Aroclor PCBs				-	
Organochlorine Pesticides	X	X	X	-	
Semivolatile Organics (SIM)	X	X	X	-	
Metals	X	-	-	X	
Mercury		-	-		
Methylmercury		-	-		
Cyanide	Х	-	X	-	
PCDD/PCDFs	X	X	X	-	
PCB Congeners	Х	X	Х	-	
Chlorinated Herbicides	Х	X	Х	-	
TOC/POC/DOC	Х	X	Х	-	
ТЕРН		-	X	-	
TSS			X		
TDS	Х		X		
Grain Size		-	-	-	

^{- =} analysis was not performed for this analytical group

x = data qualified during validation for this analytical group

3.3 Overall Accuracy/Bias

Accuracy is a measure of the bias and precision in a system, and is defined as the agreement between a measurement and an accepted reference or true value. Pre-mobil ization performance evaluation samples were analyzed prior to initiating field work. Documentation of successful analysis of the performance evaluation samples was provided to the United States Environmental Protection Agency (USEPA) by Tierra Solutions, Inc, in letters dated May 25 and October 31, 2012. Accuracy was monitored during the CSO/SWO Investigation program through the analysis of MSs, surrogate spikes, and laboratory control samples (LCSs) (performed at regular, specified intervals).

As outlined in the QAPP, the analysis of MS samples and LCSs provide laboratory results that may be compared to their associated known values to monitor potential bias. The MS and surrogate spike evaluations were used to assess bias by monitoring the actual recovery of a known quantity of a chemical, added to the native sample, versus the expected recovery. The LCS evaluations were used to assess bias by monitoring the actual recovery of a known quantity of a chemical, added to a blank, versus the expected recovery.

Acceptance criteria for each of the Accuracy evaluations described above are provided in Worksheet #12 of the QAPP. Conformance to laboratory QC sample frequency requirements, as well as acceptability of QC results for accuracy, were evaluated and considered during data verification/validation.

Data for several analytical groups associated with multiple sam pling techniques was determined to be unusable due to severe accuracy/bias issues. The following table summarizes the Phase I overall accuracy/bias quality evaluation by analytical group and sampling technique. The "x" designation indicates that an issue was identified however, such issue does not infer that the data is unusable. A more detailed discussion of this data quality parameter evaluation is provided in Section 4.1 of this report.

Overall Accuracy/Bias Issues						
Analytical Groups	Whole Water	LSM	HSM	Grab Water		
Semivolatile Organics	X	X	X	•		
Volatile Organics (trace)		-	X	-		
Aroclor PCBs	X		X	ı		
Organochlorine Pesticides	X	X	X	ı		
Semivolatile Organics (SIM)	X	X	X	-		
Metals		-	-			
Mercury		-	-			
Methylmercury		-	-			
Cyanide		-	X	-		
PCDD/PCDFs	X	X	X	-		
PCB Congeners	Х	X	X	-		
Chlorinated Herbicides	Х	X	X	-		
TOC/POC/DOC			X	-		
ТЕРН	Х	-	X	-		
TSS						
TDS						
Grain Size		-	-	-		

- = analysis was not performed for this analytical group
- x = data qualified during validation for this analytical group

3.4 Sensitivity

Sensitivity is related to the ability to compare analytical res ults with project quantitation limits (PQLs). Analytical detection limits should be at or below the PQLs to a llow effective comparisons. All sample analytical results reported during Phase I of the CSO/SWO Inves tigation were evaluated to determine if adequate sensitivity was achieved. The results for each analyt—e were cross-checked against the PQLs presented in Worksheet #15 of the QAPP. The tables in Section—3.4.1 below summarize the percent of sample results that did not meet the data quality objectives as—defined by the QAPP. The percentages expressed in these tables indicate the fraction of the total number of results reported for each analytical group and sampling technique where reporting limits exceeded the PQLs.

With respect to the results of the Phase I CSO/SWO Investigation data, there are no limitations on the data usage based on sensitivity acceptance criteria. A more detailed discussion of this data quality parameter evaluation is provided in Section 3.4.1.

3.4.1 Achieved Analytical Sensitivity

The fact that data obtained for a particular sample type/colletion technique failed to meet established PQLs for specific analytical groups as indicated in the tables below, may have impacted the number of positive results identified in those samples, thereby potentially impacting the data evaluation process. Following each table is a discussion of the analytical groups for which failure to meet the PQLs, may have impacted the Phase I data evaluation process.

Whole Water

Table 3-1
Phase1 Sensitivity Quality Evaluation for Whole Water Samples

Analytical Group	Total Number of Results Reported	Non-detected Results with PQLs Greater than those Defined in the CSO/SWO QAPP	Detected Results Between the MDL (or EDL where appropriate) and Elevated PQL	Percent of Results that did not meet Data Quality Objectives as Defined by CSO/SWO QAPP PQLs
Polychlorinated dibenzo- <i>p</i> -dioxins and dibenzofurans (PCDD/PCDFs)	102	7	42	48
Polychlorinated biphenyl (PCB) Congeners	1,008	423	77	50
Organochlorine Pesticides	112	4	8	11
Semivolatile Organics (SVOC) Selective Ion Monitoring (SIM)	120	4	4	6.7
Semivolatile Organics	200	180	7	94
Metals	92	0	7	7.6
Mercury	4	0	0	0
Methyl Mercury	4	0	0	0
VOCs	24	0	4	17
Aroclor PCBs	36	0	0	0
Chlorinated Herbicides	24	0	2	8.3
Cyanide	4	0	0	0
TOC	4	0	0	0
Total Extractable Petroleum				
Hydrocarbons (TEPH)	4	0	0	0
TSS	4	0	0	0
TDS	4	0	0	0

Each analyte group was further evaluated to determine when and if the failure to meet the PQLs may have impacted the number of positive results used to determine the r ecommended sample collection method during the Phase I evaluation process. For all analytical group s, the detected results between the method detection limit/estimated detection limit (MDL/EDL) and the ele vated PQL were included as positive results when determining the recommended sample collection method. Therefore, although the established PQLs were not met in those cases, there is no impact to the outcome of the data evaluation process.

CSO/SWO Phase I Data Quality Usability Assessment Report – Rev 2

June 2016

For the whole water (WW) PCDD/PCDF results, PQLs identified in Table 3-1 above as greater than those defined in the QAPP, all seven non-detected results were obtained from Event #1, Attempt #1, which was not included in the sample evaluation process. Therefore there was no impact on the recommended sample collection method determination.

For the WW PCB Congener results, PQLs identified in Table 3-1 bove as greater than those defined in the QAPP were only marginally exceeded due to either sample dilution prior to analyses or slightly less than targeted sample volume used for analysis. A total of 258 non-dtected results were reported above the PQL for Event #2, Attempt #2 and Event #1, Attempt #3, 20 of which were contaminants of potential concern/contaminants of potential ecological concern (COPCs/COPECs). Detection of COPCs/COPECs is prioritized when determining the recommended sample collection method, therefore these non-detected results may have impacted the number of positive COPCs/COPECs r esults identified, and could have affected the selection of a sample collection method. The remaining non-detected results reported above the PQL were obtained from Event #1, Attempt #1 and were not included in the evaluation process.

For the WW Organochlorine Pesticide results, PQLs identified in Table 3-1 above as greater than those defined in the QAPP were only marginally exceeded due to either—sample dilution prior to analyses or slightly less than targeted sample volume used for analysis. A—total of four non-detected results were reported above the PQL, all from Event #1, Attempt #2. None of these non-detected results were COPCs/COPECs, further, had the four results been positive it would not have made a significant difference in the total number of positive analytes detected. Therefore, the non-detected results did not influence the selection of a sample collection method.

For the WW SVOC SIM results, PQLs identified in Table 3-1 above as greater than those defined in the QAPP were marginally exceeded due to sample dilution prior to a nalysis. A total of four non-detected results were reported above the PQL for Event #1, Attempt #2 and Event #2, Attempt #2. Had the four results been positive it would not have made a significant difference in the total number of positive results reported (COPCs/COPECs or otherwise) and therefore the selection of a sample collection method was not impacted.

For the WW SVOC results, PQLs identified in Table 3-1 above as greater than those defined in the QAPP were exceeded to varying degrees, due to either sample dilutionprior to analysis, or use of less than targeted sample volume for analysis. A total of 90 non-detected resultswere reported above the PQL due to sample dilution for Event #1, Attempt #2. Samples collected during this event were analyzed at a dilution which resulted in a significant increase in the PQL obtained for these samples, this may have impacted the number of positive results detected, and therefore may have affected the selection of a sample collection method. The 90 non-detected SVOC results that were only marginally above the PQL due to sample volume used during the analyses for Event #2, Attempt #2, did not likely impact the number of positive results reported for that event, and therefore did not affect the selection of a sample collection method.

Low Solids Mass Dissolved

Table 3-2
Phase 1 Sensitivity Quality Evaluation for Low Solids Mass Dissolved Samples

Analytical Group	Total Number of Results Reported	Non-detected Results with PQLs Greater than those Defined in the CSO/SWO QAPP	Detected Results Between the MDL (or EDL where appropriate) and Elevated PQL	Percent of Results that did not meet Data Quality Objectives as Defined by CSO/SWO QAPP PQLs
PCDD/PCDFs	102	0	22	22
PCB Congeners	1,008	453	154	60
Organochlorine Pesticides	112	9	13	20
Semivolatile Organics SIM	120	19	18	31
Semivolatile Organics	200	7	8	7.5
Aroclor PCB	36	0	0	0
Chlorinated Herbicide	24	0	1	4.2
TOC/DOC/POC	4	0	0	0
TSS	6	1	0	17
TDS	6	0	0	0

Each analyte group was further evaluated to determine when and if the failure to meet the PQLs may have impacted the number of positive results used to determine the recommended sample collection method during the Phase I evaluation process. For all analytical groups the detected results between the MDL/EDL and the elevated PQL were included as positive results when determining the recommended sample collection method. Therefore, although the established PQLs were not met in those cases, there is no impact to the outcome of the data evaluation process.

For the low solids mass (LSM) di ssolved PCB Congener results, P QLs identified in Table 3-2 above as greater than those defined in the QAPP were only marginally exceeded due to either sample dilution prior to analyses or slightly less than targeted sample volume used f or analysis. A total of 269 non-detected results were reported above the PQL for Event #2, Attempt #2 and Event #1, Attempt #3, 24 of which were COPCs/COPECs. Detection of COPCs/COPECs is prioritized when deermining the recommended sample collection method. Therefore, these non-detected results may h ave impacted the number of positive COPC/COPECs results identified and could have affected the selection of a sample collection method. The remaining non-detected results reported above the PQL were obtained from Event #1, Attempt #1 and were not used in the sample collection evaluation process.

For the LSM dissolved Organochlorine Pesticide results, PQLs identified in Table 3-2 above as greater than those defined in the QAPP were only marginally exceeded due toeither sample dilution prior to analysis or slightly less than targeted sample volume used for analysis. A total of nine non-detected results were reported above the PQL for Event #1, Attempt #2 and Event #2, A ttempt #2, none of these non-detected results were COPCs/COPECs. Further, had those nine results been positive, it would not have made a significant difference in the total number of positive results identified. Therefore, the non-detected results did not influence the selection of a sample collection method.

For the LSM dissolved SVOC SIM results, PQLs identified in Tabl 3-2 above as greater than those defined in the QAPP were only marginally exceeded due to either sample dilution prior to analysis or less than targeted sample volume used for analysis. A total of 18 non-detected results were reported above the PQL for Event #1, Attempt #2, 10 of which were COPCs/COPECs. Detection of COPCs/COPECs is prioritized when determining the recommended sample collection method, ther efore these non-detected results may have impacted the number of pos itive COPCs/COPECs results ident ified and could have affected the selection of a sample collection method. The non-detected result reported above the PQL for Event #2, Attempt #2 was not a COPC/COPEC, further had it been positive, it would not have made a significant difference in the total number of positive results reported.

Therefore the selection of a sample collection method was not influenced in this case.

For the LSM dissolved SVOC results, PQLs identified in Table 3-2 above as greater than those defined in the QAPP were only marginally exceeded due to a less than targeted sample volume used for analysis. The seven non-detected SVOC results that were only slightly above the PQL for Event #2, Attempt #2, did not likely impact the number of positive results reported for that event, and therefore did not affect the selection of a sample collection method.

For the LSM dissolved TSS results, the PQL identified in Table 3-2 above as greater than that defined in the QAPP, has no impact on the recommended sample collection me thod determination, since TSS measurements are not used in the sample collection evaluation process.

Low Solids Mass Particulate

Table 3-3
Phase 1 Sensitivity Quality Evaluation for Low Solids Mass Particulate Samples

Analytical Group	Total Number of Results Reported	Non-detected Results with PQLs Greater than those Defined in the CSO/SWO QAPP	Detected Results Between the MDL (or EDL where appropriate) and Elevated PQL	Percent of Results that did not meet Data Quality Objectives as Defined by CSO/SWO QAPP PQLs
PCDD/PCDFs	102	0	56	55
PCB Congeners	1,008	337	155	49
Organochlorine Pesticides	112	34	13	42
Semivolatile Organic SIM	120	23	8	26
Semivolatile Organics	200	97	3	50
Aroclor PCBs	36	18	0	50
Chlorinated Herbicides	24	16	0	67
TOC/DOC/POC	4	0	0	0

Each analyte group was further evaluated to determine when and if the failure to meet the PQLs may have impacted the number of positive results used to determine the recommended sample collection method during the Phase I evaluation process. For all analytical groups the detected results between the MDL/EDL and the elevated PQL were included as positive results when determining the recommended sample collection method. Therefore, although the established PQLs were not met in those cases, there is no impact to the outcome of the data evaluation process.

CSO/SWO Phase I Data Quality Usability Assessment Report – Rev 2

June 2016

For the LSM particulate PCB Congener results, PQLs identified in Table 3-3 above as greater than those defined in the QAPP were exceeded due to both sample dilution prior to analysis and significantly less than targeted sample mass available for analysis. A total of 261 no n-detected results were reported above the PQL for Event #2, Attempt #2 and Event #1, Attempt #3, with 14of the 261 non-detected results consisting of COPCs/COPECs. Detection of C OPCs/COPECs is prioritized when determining the recommended sample collection method, therefore these non-detected results may have impacted the number of positive COPCs/COPECs results identified, and could have affected the se lection of a sample collection method. The remaining samples exhibiting non-detected results reported above the PQL were obtained from Event #1, Attempt #1 and were not included in the sample collection method evaluation process.

For the LSM particulate Organochl orine Pesticide results, PQLs identified in Table 3-3 above as greater than those defined in the QAPP were exceeded due to sample dilution prior to analysis and/or significantly less than targeted sample mass available for analysis. A total of 34 non-detected results were reported above the PQL for Event #1, Attempt #2 and Event #2, Attempt #2. If the 34 results had been positive it may have made a significant difference in the total number of **p**sitive results identified and therefore could have had an impact on the selection of a sample collection method.

For the LSM particulate SVOC SIM results, PQLs identified in Ta ble 3-3 above as greater than those defined in the QAPP were exceeded due to sample dilution prior to analysis and/or significantly less than targeted sample mass available for analysis. A total of 18 nondetected results were reported above the PQL for Event #1, Attempt #2, in which nine were COPCs/COPECs. Detetion of COPCs/COPECs is prioritized when determining the recommended sample collection method, ther efore these non-detected results may have impacted the number of positive COPCs/COPECs results identified, and could have affected the selection of a sample collection method. The five non-detected results reported above the PQL for Event #2, Attempt #2 were not COPC/COPECs, Further, had they been positive it would not have made a significant difference in the total number of positive results reported. Therefore, the selection of a sample collection method was not influenced in this case.

For the LSM particulate SVOC results, PQLs identified in Table3-3 above as greater than those defined in the QAPP were exceeded due to significantly less than targetedsample mass available for analysis. A total of 97 non-detected results were reported above the PQL all from Event #2, Attempt #2. Had the 97 results been positive it may have made a significant difference in the total number of positive results identified and therefore could have had an impact on the selection of a sample collection method.

For the LSM particulate Aroclor PCB results, PQLs identified in Table 3-3 above as greater than those defined in the QAPP were exceeded due to significantly less that a natural natur

For the LSM particulate Herbicide results, PQLs identified in Table 3-3 above as greater than those defined in the QAPP were exceeded due to significantly less than target ed sample mass available for analysis. A total of 16 non-detected results were reported above the PQL fo r Event #2, Attempt #2 and Event #1, Attempt #3. Had the 16 results been positive it may have madea significant difference in the total number of positive results identified and therefore could have had an impact on the selection of a sample collection method.

High Solids Mass Dissolved

Table 3-4
Phase 1 Sensitivity Quality Evaluation for High Solids Mass Dissolved Samples

Analytical Group	Total Number of Results Reported	Non-detected Results with PQLs Greater than those Defined in the CSO/SWO QAPP	Detected Results Between the MDL (or EDL where appropriate) and Elevated PQL	Percent of Results that did not meet Data Quality Objectives as Defined by CSO/SWO QAPP PQLs
PCDD/PCDFs	102	0	48	47
PCB Congeners	1,008	446	128	57
Organochlorine Pesticides	112	4	18	20
Semivolatile Organics SIM	120	0	6	5.0
Semivolatile Organics	200	140	7	74
VOCs	24	0	4	17
Aroclor PCBs	36	0	0	0
Chlorinated Herbicides	24	3	3	25
Cyanide	4	0	0	0
TOC	4	0	0	0
ТЕРН	4	0	0	0
TSS	8	0	0	0
TDS	8	0	0	0

Each analyte group was further evaluated to determine when and if the failure to meet the PQLs may have impacted the number of positive results used to determine the recommended sample collection method during the Phase I evaluation process. For all analytical groups the detected results between the MDL/EDL and the elevated PQL were included as positive results when determining the recommended sample collection method. Therefore, although the established PQLs were not met in those cases, there is no impact to the outcome of the data evaluation process.

For the high solids mass (HSM) dissolved PCB Congener results, PQLs identified in Table 3-4 above as greater than those defined in the QAPP were exceeded due to sample dilution prior to analysis and/or use of slightly less than targeted sample volume for analysis. A otal of 293 non-detected results were reported above the PQL for Event #2, Attempt #2 and Event #1, Attempt #3, 23 of which were COPCs/COPECs. Detection of COPCs/COPECs is prioritized when determining the commended sample collection method, therefore these non-detected results may have impacted the numb er of positive COPCs/COPECs results identified, and could have affected the selection of a sample ollection method. The remaining non-detected results reported above the PQL were obtained from Event #1, Att empt #1 and were not included in the evaluation process.

For the HSM dissolved Organochlorine Pesticide results, PQLs id entified in Table 3-4 above as greater than those defined in the QAPP were exceeded due to sample diltion prior to analysis and/or use of slightly less than targeted sample volume for analysis. A total of fouron-detected results were reported above the PQL for Event #1, Attempt #2 and Event #2, Attempt #2, none of which were COPCs/COPECs. Further, had the four results been positive it would not have made a sig nificant difference in the total number of positive analytes detected. Therefore, the non-detected results did not influence the selection of a sample collection method.

For the HSM dissolved SVOC results, PQLs identified in Table 3-4 above as greater than those defined in the QAPP were exceeded due to sample dilution prior to analysis and/or use of slightly less than targeted sample volume for analysis. The 51 non-detected results reported above the PQL for Event #1, Attempt #2 did not affect the selection of the sample collection method, a both the primary and duplicate samples were eliminated from consideration because more than ten percent of the results reported were rejected during data validation. A total of 89 non-detected results were reported above the PQL for Event #2, Attempt #2.

For the HSM dissolved Herbicide results, PQLs identified in Table 3-4 above as greater than those defined in the QAPP were exceeded due to use of less than targeted samp le volume for analysis. A total of three non-detected results were reported above the PQL all from Even#1, Attempt #2. Had the three results been positive it may have made a significant difference in the total number of positive results identified and therefore could have had an impact on the selection of a sample collection method.

High Solids Mass Particulate

Table 3-5
Phase 1 Sensitivity Quality Evaluation for High Solids Mass Particulate Samples

Analytical Group	Total Number of Results Reported	Non-detected Results with PQLs Greater than those Defined in the CSO/SWO QAPP	Detected Results Between the MDL (or EDL where appropriate) and Elevated PQL	Percent of Results That Did Not Meet Data Quality Objectives as Defined by CSO/SWO QAPP PQLs
PCDD/PCDFs	102	5	12	17
PCB Congeners	1,008	308	79	38
Organochlorine Pesticides	112	38	10	43
SVOC SIM	120	13	1	12
SVOC	200	178	10	94
VOCs	42	28	11	93
Aroclor PCBs	36	26	5	86
Chlorinated Herbicides	24	0	16	67
Cyanide	6	0	0	0
TOC	6	0	0	0
ТЕРН	4	0	0	0

EDL = estimated detection limit

MDL = method detection limit

Each analyte group was further evaluated to determine when and if the failure to meet the PQLs may have impacted the number of positive results used to determine the recommended sample collection method during the Phase I evaluation process. For all analytical groups the detected results between the MDL/EDL

CSO/SWO Phase I Data Quality Usability Assessment Report – Rev 2

June 2016

and the elevated PQL were included as positive results when det ermining the recommended sample collection method. Therefore, although the established PQLs were not met in those cases, there is no impact to the outcome of the data evaluation process.

For the HSM particulate PCDD/PCDFs results, PQLs identified in Table 3-5 above as greater than those defined in the QAPP were exceeded due to sample dilution prior—to analysis, and/or less than targeted sample mass used for analysis. A total of three non-detected sults were reported above the PQL for Event #2, Attempt #2. Since a significantly greater number of positive COPCs/COPECs were already identified in the HSM sample than others, had the three results been posit ive it would not have made a significant difference in the selection of a sample collection method. One non-detected result for Event #1, Attempt #3 was a COPC/COPEC. Detection of COPCs/COPECs is prioritized when determining the recommended sample collection method, therefore this non-detected result may have impacted the number of positive COPCs/COPECs results identified, and could have affected the selection of a sample collection method.

For the HSM particulate PCB Congener results, PQLs identified in Table 3-5 above as greater than those defined in the QAPP were exceeded due to sample dilution prior to analysis, and/or less than targeted sample mass used for analysis. A total of 212 non-detected results were reported above the PQL for Event #2, Attempt #2 and Event #1, Attempt #3, nine of which were COP Cs/COPECs. Since a significantly greater number of positive COPCs/COPECs were already identified in the HSM sample than others, had the nine results been positive it would not have made a significant difference in the selection of a sample collection method. The remaining non-detected results reported above the PQL were obtained from Event #1, Attempt #1 and were not included in the sample collection method evaluation process.

For the HSM particulate Organochlorine Pesticide results, PQLs identified in Table 3-5 above as greater than those defined in the QAPP were exceeded due to sample dilu tion prior to analysis, and/or less than targeted sample mass used for analysis. A total of 38 non-dete cted results were reported above the PQL for Event #1, Attempt #2 and Event #2, Attempt #2, none of whic h were COPCs/COPECs. Since a significantly greater number of positive COPCs/COPECs were already identified in the HSM sample than others, had the 38 results been positive it would not have madea significant difference in the selection of a sample collection method.

For the HSM particulate SVOC SIM results, PQLs identified in T able 3-5 above as greater than those defined in the QAPP were marginally exceeded due to sample dilution prior to analysis, less than targeted sample mass used for analysis and/or the percent solids of the samples. A total of 13 non-detected results were reported above the PQL for Event #1, Attempt #2 and Event #2, Attempt #2, five of which were COPCs/COPECs. Detection of COPCs/COPECs is prioritized when determining the recommended sample collection method, therefore these non-detected results may have impacted the number of positive COPCs/COPECs results identified, and could have affected the selection of a sample collection method.

For the HSM particulate SVOC results, PQLs identified in Table3-5 above as greater than those defined in the QAPP were exceeded due to sample dilution prior to analysis, less than targeted sample mass used for analysis and/or the percent solids of the samples. A total of 86 non-detected results were reported above the PQL for Event #2, Attempt #2. Had the 86 results been posit ive it may have made a significant difference in the total number of positive results identified and therefore could have had an impact on the selection of a sample collection method. Quality control issues identified in the primary and duplicate analyses of Event #1, Attempt #2, HSM dissolved analyses eliminated the HSM sample collection method from consideration, resulting in an inconclusive overall determination for that Event/Attempt. Therefore

CSO/SWO Phase I Data Quality Usability Assessment Report – Rev 2

June 2016

the 92 PQLs exceeded with non-detected results in the HSM particulate component of Event #1, Attempt #2, would have had no impact on selection of a sample collection method.

For the HSM particulate VOC results, PQLs identified in Table 3-5 above as greater than those defined in the QAPP were marginally exceeded due to less than targeted sample mass available for analysis and/or the percent solids of the samples. The non-detected results reported above the PQL for Event #1, Attempt #2 and Event #2, Attempt #1, did not affect the selection of a sample collection method as the high solids mass samples had a significant amount of rejected data (see Section 4.1.6 for a description of rejected data), and were eliminated from consideration on that basis.

For the HSM particulate Aroclor PCB results, PQLs identified in Table 3-5 above as greater than those defined in the QAPP were exceeded due to the percent solids of the samples. A total of seven non-detected results were reported above the PQL for Event #2, Attempt #2 (o riginal sample), all of which were COPCs/COPECs. Detection of COPCs/COPECs is prioritized when determining the recommended sample collection method, therefore these non-detected results may have impacted the number of positive COPCs/COPECs results identified, and could have affected the se lection of a sample collection method. The 19 non-detected results above the PQL for Event #1, Attempt #2 and Event #2, Attempt #2 (field duplicate only) did not likely impact the selection of a sample collection method, since a larger number of positive COPC/COPECs were already identified in the HSM sample collected during these events than other sample collection methods.

3.5 Representativeness

Representativeness is the degree to which a data set accurately presents the characteristics of a population, parameter conditions at a sample point, or an environmental con dition. Data are representative when all sampling and analyses are performed in compliance with appropri ate procedures. Performing sample analyses within the specified holding times and adhering to sample handling and storage requirements are also critical elements in obtaining representative sample data. These elements were evaluated and considered during data verification/validation. Acceptance cri teria for sample handling, storage and holding times are provided in Worksheets #19-1 of the QAPP.

With respect to the results of the Phase I CSO/SWO Investigation data, there are no limitations on the data usage based on representativeness acceptance criteria. The following table summarizes the Phase I representativeness quality evaluation by analytical group and sampling technique. The "x" designation indicates that an issue was identified however, such issue does not infer that the data is unusable. A more detailed discussion of this data quality parameter evaluation is provided in Section 4.1 of this report.

Holding Time Violations					
Analytical Groups	Whole Water	LSM	MSH	Grab Water	
Semivolatile Organics			X	-	
Volatile Organics (trace)		-		•	
Aroclor PCBs	X	X		•	
Organochlorine Pesticides			X	ı	
Semivolatile Organics (SIM)	X		X	ı	
Metals		-	ı		
Mercury	X	-	ı		
Methylmercury		-	ı		
Cyanide		-		ı	
PCDD/PCDFs		X		ı	
PCB Congeners				-	
Chlorinated Herbicides		X		-	
TOC/POC/DOC		X		-	
ТЕРН		-	X	-	
TSS				X	
TDS				X	
Grain Size		-	-	-	

^{- =} analysis was not performed for this analytical group

3.6 Comparability

Comparability expresses the confidence with which one set of data can be compared to another to measure the same property. Data can be compared to the degree that their accuracy, precision, and representativeness are known and documented. Data are comparable if QC measures such as collection techniques, measurement procedures, analytical methods, and reporting units are equivalent for the samples within a sample set. Data subject to established quality assur ance/quality control (QA/QC) measures are deemed more reliable and, therefore, more comparable, than data generated without such measures.

Consistent application of prescribed procedures was monitored t hroughout Phase I of the CSO/SWO Investigation program. Likewise, specific data verification/validation protocols were consistently applied to all data generated under this program to understand and document accuracy/bias, accuracy/bias contamination, precision, sensitivity and representativeness, thereby establishing comparability as defined above.

During data validation activities, analytical data were evaluated using a defined set of guidelines and acceptance criteria. In addition, data validation qualifiers were consistently applied to the analytical data generated during the Phase I CSO/SWO Investigation program. The data validation process serves to increase the degree of data comparability achieved.

x = data qualified during validation for this analytical group

With respect to the results of the Phase I CSO/SWO Investigation data, there are no limitations on the data usage based on representativeness acceptance criteria.

3.7 Field and Analytical Completeness

There are two measures of completeness defined for the CSO/SWO Investigation program: field completeness and analytical completeness. Field completeness is defined as the ratio of the number of samples received in acceptable condition by the laboratories to the number of samples planned to be collected as specified in the QAPP. Analytical completeness is defined as the ratio of total analytical data results reported to the total number of analytical results requested on samples submitted for analysis. The formulas used to compute field and analytical completeness are presented below.

The targeted field and analytical completeness goals were 90% for the CSO/SWO Investigation program; these goals were met, or exceeded, as summarized below.

CSO/SWO Investigation	Completeness Goal Established in CSO/SWO Investigation QAPP	Phase I CSO/SWO Investigation Completeness Achieved
Field Completeness (Overall)	90%	100%
Analytical Completeness (Overall)	90%	100%

Phase I CSO/SWO Investigation Field Completeness by Analysis and Collection Method

	Number o	of Samples Co	ollected by Sa	ımple Type	Total Number of	Total Number	Completeness
Analytical Group	Whole Water	LSM ¹	HSM ¹	Grab Water ²	Samples Collected	of Samples Planned	Achieved (%)
Semivolatile Organics	4	8	8	-	20	20	100
Volatile Organics	4	-	8	-	12	12	100
Aroclor PCBs	4	8	8		20	20	100
Organochlorine Pesticides	4	8	8	_	20	20	100
Semivolatile Organics (SIM)	4	8	8	_	20	20	100
Metals	4	-	-	8	12	12	100
Mercury	4	-	-	8	12	12	100
Methylmercury	4	-	-	8	12	12	100
Cyanide	4	-	8	-	12	12	100
PCDD/PCDFs	4	8	8	-	20	20	100
PCB Congeners	4	8	8	-	20	20	100
Chlorinated Herbicides	6	12	12	-	30	20	150
TOC/POC/DOC ³	4	8	8	-	20	20	100
TEPH	4	-	8	-	12	12	100
TSS	6	6	6	-	18	12	150
TDS	6	6	6	-	18	12	150
Grain Size	4	-	-	-	4	4	100

^{1 –} Particulate and dissolved samples

Phase 1 CSO/SWO Investigation Analytical Completeness by Analysis and Collection Method Whole Water

Analytical Group	Samples Analyzed Including Trip Blanks	Analytes per Sample	Total Results	Rejected Results	Analytical Completeness Achieved
Semivolatile Organics	4	50	200	0	100%
Volatile Organics	6	6	36	0	100%
Aroclor PCBs	4	9	36	0	100%
Organochlorine Pesticides	4	28	112	0	100%
Semivolatile Organics (SIM)	4	30	120	0	100%
Metals	3	23	69	0	100%
Mercury	3	1	3	0	100%
Methyl mercury	3	1	3	0	100%
Cyanide	4	1	4	0	100%
PCDD/PCDFs	6	17	102	0	100%

^{2 –} Total and dissolved samples

^{3 –} TOC, POC and DOC analyses are mutually exclusive. Therefore, only one of the three analyses is performed per sample type.

June 2016

PCB Congeners	6	168	1008	0	100%
Chlorinated Herbicides	6	4	24	0	100%
TOC	4	1	4	0	100%
ТЕРН	4	1	4	0	100%
Grain Size	4	85	340	0	100%
TSS	6	1	6	0	100%
TDS	6	1	6	0	100%

LSM Particulate

Analytical Group	Samples Analyzed Including Trip Blanks	Analytes per Sample	Total Results	Rejected Results	Analytical Completeness Achieved
Semivolatile Organics	4	50	200	10	95%
Aroclor PCBs	4	9	36	0	100%
Organochlorine Pesticides	4	28	112	1	99%
Semivolatile Organics (SIM)	4	30	120	0	100%
PCDD/PCDFs	6	17	102	0	100%
PCB Congeners	6	168	1008	0	100%
Chlorinated Herbicides	6	4	24	0	100%
POC	4	1	4	0	100%

LSM Dissolved

Analytical Group	Samples Analyzed Including Trip Blanks	Analytes per Sample	Total Results	Rejected Results	Analytical Completeness Achieved
Semivolatile Organics	4	50	200	1	99.5%
Aroclor PCBs	4	9	36	0	100%
Organochlorine Pesticides	4	28	112	0	100%
Semivolatile Organics (SIM)	4	30	120	0	100%
PCDD/PCDFs	6	17	102	0	100%
PCB Congeners	6	168	1008	0	100%
Chlorinated Herbicides	6	4	24	0	100%
DOC	4	1	4	0	100%
TSS	4	1	4	0	100%
TDS	4	1	4	0	100%

HSM Particulate

Analytical Group	Samples Analyzed Including Trip Blanks	Analytes per Sample	Total Results	Rejected Results	Analytical Completeness Achieved
Semivolatile Organics	4	50	200	2	99%
Volatile Organics	9	6	54	25	53.7%
Aroclor PCBs	4	9	36	0	100%
Organochlorine Pesticides	4	28	112	6	94.6%
Semivolatile Organics (SIM)	4	30	120	0	100%
Cyanide	4	1	4	0	100%
PCDD/PCDFs	6	17	102	0	100%
PCB Congeners	6	168	1008	0	100%
Chlorinated Herbicides	6	4	24	0	100%
TOC	4	1	4	0	100%
ТЕРН	4	1	4	0	100%

HSM Dissolved

Analytical Group	Samples Analyzed Including Trip Blanks	Analytes per Sample	Total Results	Rejected Results	Analytical Completeness Achieved
Semivolatile Organics	4	50	200	16	92%
Volatile Organics	6	6	36	0	100%
Aroclor PCBs	4	9	36	0	100%
Organochlorine Pesticides	4	28	112	0	100%
Semivolatile Organics (SIM)	4	30	120	0	100%
Cyanide	4	1	4	0	100%
PCDD/PCDFs	6	17	102	0	100%
PCB Congeners	6	168	1008	0	100%
Chlorinated Herbicides	6	4	24	0	100%
DOC	4	1	4	0	100%
ТЕРН	4	1	4	0	100%
TSS	8	1	8	0	100%
TDS	8	1	8	0	100%

Grab Samples

Analytical Group	Samples Analyzed Including Trip Blanks	Analytes per Sample	Total Results	Rejected Results	Analytical Completeness Achieved
Metals	12	23	276	0	99.5%
Mercury	12	1	12	0	100%
Methylmercury	12	1	12	0	100%
TSS	2	1	2	0	100%
TDS	1	1	1	0	100%

4. Phase I CSO/SWO Investigation Data Verification/Validation

Phase I CSO/SWO Investigation analytical results were provided by the laboratories both electronically and in hard copy format. Upon receipt from the laboratory, results for specific analytical groups described below were verified or validated by Environmental Data Services, Ltd. (EDS) using the following procedures:

Semivolatile Organics	USEPA Region 2 SOP HW-35, Revision 1
Volatile Organics (trace)	USEPA Region 2 SOP HW-34, Revision 1
Aroclor PCBs	USEPA Region 2 SOP HW-37, Revision 1
Organochlorine Pesticides	EDS SOP: Organochlorine Pesticides by HRGC/HRMS USEPA 1699, Rev .0, 7/10
Semivolatile Organics (SIM)	USEPA Region 2 HW-35, Revision 1
Metals	EDS SOP: Metals by ICP/MS USEPA 1638, Rev.0, 7/10
Mercury	EDS SOP: Mercury by CVAFS USEPA 1631, Rev.0, 7/10
Methylmercury	EDS SOP: Methyl Mercury by CVAFS USEPA 1630, Rev.0, 7/10
Cyanide	USEPA Region 2 SOP HW-2, Revision 13
PCDD/PCDFs	USEPA Region 2 SOP HW-25, Revision 3
PCB Congeners	EDS SOP: Congener PCB, Rev. 3, 7/10
Chlorinated Herbicides	USEPA Region 2 SOP HW-17, Revision 3
TOC (solid/liquid)/DOC/POC	EDS SOP:TOC-01 Rev.2, 7/10
ТЕРН	EDS SOP:TEPH-01 Rev. 3, 7/07
TSS	EDS SOP: TSS by Gravimetric SM 2540D, Rev. 0, 7/10
TDS	EDS SOP: TDS by Gravimetric SM 2540C, Rev. 0, 7/10
Grain Size	SOP-14, Revision 2 – Verification/Validation Geotechnical Data

The verification/validation standard operating procedures (SOPs), as referenced above, are provided in Appendix C of the QAPP. The data verification/validation process is detailed in Worksheets #34, 35, and 36 of the QAPP.

4.1 Data Quality Issues

Two types of data quality issues are discussed in this section; systematic data quality issues and random data quality issues. Systematic data quality issues are thosehat are identified as having a consistent impact on the quality of results reported (i.e., data quality of all samples and/or analytical groups are affected by a single data quality issue), due to a common circumstance or proedural application. Systematic data quality issues are described in Sections 4.1.1, 4.1.3, 4.1.5, and 4.1.7 as well as incorporated into Sections 4.1.2, 4.1.4, 4.1.6, and 4.1.8. Random data quality issues are thosehat do not have a consistent impact the quality of results (i.e., data quality for a specific sample(s) and/oranalyte(s) are affected by the data quality issue). Random data quality issues are presented in Sections 4.1.2, 4.1.4, 4.1.6, and 4.1.8.

Sections 4.1.2, 4.1.4, 4.1.6, and 4.1.8 summarizes the data val idation findings related to systematic and random data quality issues for each analytical group. These validation findings have been separated into two distinct categories, major data quality issues and minor data quality issues. Major data quality issues are those that result in the qualification of the analytical value reported as "R", or rejected. This occurs due to the presence of significant QA/QC problems that render the analysis invalid and the results unusable. Minor data quality issues include all other QA/QC problems identified during the data validation process that require sample results to be qualified, indicating some level of uncertainty associated with the reported result. Qualifiers applied to sample results were assigned based on the validation protocols specified in Worksheet #36 of the QAPP.

Conclusions based on the information presented in these summaries can be found in Section 5 of this report.

4.1.1 Whole Water Samples Systematic Data Quality Issues

Four systematic data quality issues were identified during the Phase I CSO/SWO Investigation data whole water sample validation task. These systematic data quality issues are summarized below:

All internal standard recoveries for 13C-PCB-205 were outside the quality control limits. All results for PCB-205 were qualified as estimated.
All field blanks contained hexachlorobenzene, 2,4'-DDE, 4,4'-DDE, 2,4'-DDD, 2,4'-DDT, 4,4'-DDD and 4,4'-DDT resulting in the positive results being qualified non-detected "U".
All field blanks contained butylbenzylphthalate resulting in the positive results being qualified non-detected "U".
All surrogate recoveries for Decachlorobiphenyl were outside the quality control limit. All non-detected results for Aroclors were qualified as estimated.

4.1.2 Whole Water Samples Systematic and Random Data Quality Issues by Analytical Group

Semivolatile Organic Compounds

The Phase I CSO/SWO Investigation whole water sample SVOC dataset is comprised of four samples with 200 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation SVOC analyses.

Five minor data quality issues were identified in the Phase I C SO/SWO Investigation whole water SVOC dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
Semivolatile Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected		
Field blank contamination	Accuracy/Bias Contamination	200	4	4	2.0		
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	200	4	6	3.0		
Non-compliant internal standard recovery	Overall Accuracy/Bias	200	1	1	0.50		
Non-compliant method surrogate recovery	Overall Accuracy/Bias	200	3	9	4.5		
Non-compliant project specific surrogate recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	200	4	14	7.0		

Volatile Organic Compounds (trace)

The Phase I CSO/SWO Investigation whole water VOC (trace) dataset is comprised of four samples with 24 associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation VOC (trace) analyses.

Aroclor Polychlorinated Biphenyls

The Phase I CSO/SWO Investigation whole water Aroclor PCB dataset is comprised of four samples with 36 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Aroclor PCB analyses.

Three minor data quality issues were identified in the Phase CSO/SWO Investigation whole water Aroclor PCB dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
Aroclor PCB Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Aroclor PCB Results Affected		
Non-compliant holding time	Representativeness	36	2	18	50.0		
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	36	2	2	5.6		
Non-compliant method surrogate recovery	Overall Accuracy/Bias	36	4	36	100		

Organochlorine Pesticides

The Phase I CSO/SWO Investigation whole water Organochlorine Pe sticide dataset is comprised of four samples with 112 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Organochlorine Pesticide analyses.

Six minor data quality issues were identified in the Phase I CSO/SWO Investigation whole water Organochlorine Pesticide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues									
Organochlorine Pesticide Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Organochlorine Pesticide Results Affected				
Field blank contamination	Accuracy/Bias Contamination	112	4	29	25.9				
Non-compliant qualitative requirements	Overall Accuracy/Bias	112	1	1	0.9				
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	112	2	6	5.4				
Non-compliant field duplicate relative percent difference	Precision	112	2	2	1.8				
Non-compliant internal standard recovery	Overall Accuracy/Bias	112	3	74	66.1				
Non-complaint project specific surrogate recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	112	4	22	19.6				

Semivolatile Organic Compounds - Selective Ion Monitoring

The Phase I CSO/SWO Investigation whole water SVOCs SIM dataset is comprised of four samples with 120 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation SVOCs SIM analyses.

Six minor data quality issues were identified in the Phase I CSO/SWO Investigation whole water SVOCs SIM dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
Semivolatile SIM Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC SIM Results Affected		
Non-compliant holding time	Representativeness	120	2	60	50.0		
Method blank contamination	Accuracy/Bias Contamination	120	2	2	1.7		
Field blank contamination	Accuracy/Bias Contamination	120	3	23	19.2		
Non-compliant initial calibration relative standard deviation	Overall Accuracy/Bias	120	2	2	1.7		
Non-compliant project specific surrogate recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	120	1	7	5.8		
Non-compliant field duplicate relative percent difference	Precision	120	4	64	53.3		

Metals

The Phase I CSO/SWO Investigation whole water Metals dataset is comprised of four samples with 92 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Metals analyses.

Two minor data quality issues were identified in the Phase I CSO/SWO Investigation whole water Metals dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Metals Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Metals Results Affected
Field blank contamination	Accuracy/Bias Contamination	92	4	6	6.5
Continuing calibration blank contamination	Accuracy/Bias Contamination	92	2	4	4.4

Mercury

The Phase I CSO/SWO Investigation whole water Mercury dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Mercury analyses.

Two minor data quality issues were identified in the Phase I CSD/SWO Investigation whole water Mercury dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Mercury Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Mercury Results Affected
Non-compliant holding time	Representativeness	4	4	4	100
Non-compliant field duplicate relative percent difference	Precision	4	4	4	100

Methyl Mercury

The Phase I CSO/SWO Investigation whole water Methyl Mercury data set is comprised of four samples with four associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation Methyl Mercury analyses.

Cyanide

The Phase I CSO/SWO Investigation whole water Cyanide dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Cyanide analyses.

One minor data quality issue was identified in the Phase I CSO/SWO Investigation whole water Cyanide dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
Cyanide Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Cyanide Results Affected
Field blank contamination	Accuracy/Bias Contamination	4	2	2	50.0

Polychlorinated Dibenzo-p-dioxins / Polychlorinated Dibenzofurans

The Phase I CSO/SWO Investigation whole water PCDD/PCDFs dataset is comprised of six samples with 102 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCDD/PCDF analyses.

Five minor data quality issues were identified in the Phase I C SO/SWO Investigation whole water PCDD/PCDF dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
PCDD/PCDFs Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCDD/PCDF Results Affected		
Field blank contamination	Accuracy/Bias Contamination	102	2	7	6.9		
Non-compliant matrix spike/matrix spike duplicate recovery	Overall Accuracy/Bias	102	1	1	1.0		
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	102	1	1	1.08		
Non-compliant field duplicate relative percent difference	Precision	102	4	10	9.8		
Non-complaint project specific labeled analog recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	102	4	17	16.7		

Polychlorinated Biphenyl Congeners

The Phase I CSO/SWO Investigation whole water PCB Congener dataset is comprised of six samples with 1,008 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCB Congener analyses.

Four minor data quality issues were identified in the Phase I C SO/SWO Investigation whole water PCB Congener dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
PCB Congeners Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCB Congener Results Affected		
Field blank contamination	Accuracy/Bias Contamination	1,008	5	123	12.2		
Non-compliant field duplicate relative percent difference	Precision	1,008	6	266	26.4		
Non-compliant internal standard recovery	Overall Accuracy/Bias	1,008	6	308	30.6		
Non-complaint project specific labeled analog recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	1,008	2	58	5.8		

Chlorinated Herbicides

The Phase I CSO/SWO Investigation whole water Chlorinated Herbi cide dataset is comprised of six samples with 24 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Chlorinated Herbicide analyses.

Four minor data quality issues were identified in the Phase I C SO/SWO Investigation whole water Chlorinated Herbicide dataset. The indentified minor data quality issues are described in the table below.

Minor Data Quality Issues							
Chlorinated Herbicide Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Chlorinated Herbicide Results Affected		
Method blank contamination	Accuracy/Bias Contamination	24	2	2	8.3		
Field blank contamination	Accuracy/Bias Contamination	24	4	7	29.2		
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	24	2	2	8.3		
Non-compliant dual column analysis percent difference	Precision	24	4	9	37.5		

Total Organic Carbon

The Phase I CSO/SWO Investigation whole water TOC dataset is co mprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TOC analyses.

One minor data quality issue was identified in the Phase I CSO/ SWO Investigation whole water TOC dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
TOC Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TOC Results Affected
Field blank contamination	Accuracy/Bias Contamination	4	2	2	50.0

Total Extractable Petroleum Hydrocarbon

The Phase I CSO/SWO Investigation whole water TEPH dataset is c omprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO TEPH Investigation analyses.

Three minor data quality issues were identified in the Phase ICSO/SWO Investigation whole water TEPH data set. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
TEPH Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TEPH Results Affected			
Non-compliant initial calibration relative standard deviation	Overall Accuracy/Bias	4	2	2	50.0			
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	4	2	2	50.0			
Non-compliant field duplicate relative percent difference	Precision	4	4	4	100			

Total Suspended Solids

The Phase I CSO/SWO Investigation whole water TSS dataset is co mprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TSS analyses.

One minor data quality issue was identified in the Phase I CSO/ SWO Investigation whole water TSS dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues							
TSS Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TSS Results Affected		
Non-compliant field duplicate relative percent difference	Precision	4	4	4	100		

Total Dissolved Solids

The Phase I CSO/SWO Investigation whole water TDS dataset is co mprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TDS analyses.

One minor data quality issue was identified in the Phase I CSO/ SWO Investigation whole water TDS dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues							
TDS Whole Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TDS Results Affected		
Field blank contamination	Accuracy/Bias Contamination	4	2	2	50.0		

Geotechnical

The Phase I CSO/SWO Investigation Whole Water grain size datase t is comprised of four samples with 340 associated results.

No major or minor data quality issues were identified during the verification of the Phase I CSO/SWO Investigation grain size analyses.

4.1.3 Low Solids Mass Samples Systematic Data Quality Issues

Four systematic data quality issue was identified during the Ph ase I CSO/SWO Investigation data LSM sample validation task. These systematic data quality issues are summarized below:

Field blanks associated with all samples contained hexachlorobenzene, 4,4'-DDE, 2,4'-DDD, 2,4'-DDT, 4,4'-DDD and 4,4'-DDT resulting in the positive results being qualified non-detected "U".
All closing continuing calibration percent differences for Di-n-octylphthalate were outside the quality control limit. All results for Di-n-octylphthalate were qualified as estimated.
All field blanks contained PCB-11, PCB-16/32, PCB-17, PCB-18, PCB-19, PCB-20/21/33 and PCB-22 resulting in the positive results being qualified non-detected "U".
Due to actual TSS values being lower than estimated, LSM Particulate sample masses were much lower than anticipated. This resulted in all analytical groups having reporting limits well in excess of project quantitation limits stated in the QAPP.

4.1.4 Low Solids Mass Samples Systematic and Random Data Quality Issues by Analytical Group

Low Solids Mass Dissolved

Semivolatile Organic Compounds

The Phase I CSO/SWO Investigation LSM dissolved sample SVOC dataset is comprised of four samples with 200 associated results.

One major data quality issue was identified during validation of the Phase I CSO/SWO Investigation LSM dissolved SVOC analyses. The internal standard perylene-d12 ex hibited a recovery below the quality control limit for sample PR1CSOCLYLD-01B foi-nd-octylphthalate. The identified major data quality issue is described in the table below.

Major Data Quality Issues							
Semivolatile LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected		
Extremely poor internal standard recovery	Overall Accuracy/Bias	200	1	1	0.50		

Five minor data quality issues were identified in the Phase I SO/SWO Investigation LSM dissolved SVOC dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
Semivolatile LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected			
Field blank contamination	Accuracy/Bias Contamination	200	3	4	2.0			
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	200	4	8	4.0			
Non-compliant method surrogate recovery	Overall Accuracy/Bias	200	2	6	3.0			
Non-compliant project specific surrogate recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	200	4	23	11.5			
Non-compliant field duplicate relative percent difference	Precision	200	2	4	2.0			

Aroclor Polychlorinated Biphenyls

The Phase I CSO/SWO Investigation LSM dissolved sample Aroclor PCB dataset is comprised of four samples with 36 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Aroclor PCB analyses.

One minor data quality issue was identified in the Phase I CSOSWO Investigation LSM dissolved Aroclor PCB dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
Aroclor PCBs LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Aroclor PCB Results Affected
Non-compliant holding time	Representativeness	36	2	18	50.0

Organochlorine Pesticides

The Phase I CSO/SWO Investigation LSM dissolved sample Organochorine Pesticide dataset is comprised of four samples with 112 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Organochlorine Pesticide analyses.

Four minor data quality issues were identified in the Phase I C SO/SWO Investigation LSM dissolved Organochlorine Pesticide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
Organochlorine Pesticides LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Organochlorine Pesticide Results Affected			
Field blank contamination	Accuracy/Bias Contamination	112	4	30	26.8			
Non-compliant internal standard recovery	Overall Accuracy/Bias	112	2	44	39.3			
Non-compliant project specific labeled analog recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	112	4	18	16.1			
Non-compliant field duplicate relative percent difference	Precision	112	2	2	1.8			

Semivolatile Organic Compounds - Select Ion Monitoring

The Phase I CSO/SWO Investigation LSM dissolved sample SVOCs SI M dataset is comprised of four samples with 120 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation SVOCs SIM analyses.

Seven minor data quality issues were identified in the Phase I CSO/SWO Investigation LSM dissolved SVOCs SIM dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
Semivolatile SIM LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC SIM Results Affected		
Field blank contamination	Accuracy/Bias Contamination	120	4	26	21.7		
Non-compliant initial calibration relative standard deviation	Overall Accuracy/Bias	120	2	2	1.7		
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	120	2	2	1.7		
Non-compliant method surrogate recovery	Overall Accuracy/Bias	120	1	16	13.3		
Non-compliant project specific surrogate recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	120	1	16	13.3		
Non-compliant matrix spike/matrix spike duplicate recovery	Overall Accuracy/Bias	120	1	12	10.0		
Non-compliant field duplicate relative percent difference	Precision	120	2	14	11.7		

Polychlorinated Dibenzo-p-dioxins / Polychlorinated Dibenzofurans

The Phase I CSO/SWO Investigation LSM dissolved sample PCDD/PCD Fs dataset is comprised of six samples with 102 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCDD/PCDFs analyses.

Three minor data quality issues were identified in the Phase I CSO/SWO Investigation LSM dissolved PCDD/PCDFs dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
PCDD/PCDFs LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCDD/PCDF Results Affected
Non-compliant holding time	Representativeness	102	2	34	33.3
Field blank contamination	Accuracy/Bias Contamination	102	4	10	9.8
Non-compliant project specific labeled analog recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	102	6	63	61.8

Polychlorinated Biphenyl Congeners

The Phase I CSO/SWO Investigation LSM dissolved PCB Congener da taset is comprised of six samples with 1,008 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCB Congener analyses.

Three minor data quality issues were identified in the Phase CSO/SWO Investigation LSM dissolved PCB Congener dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
PCB Congeners LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCB Congener Results Affected		
Method blank contamination	Accuracy/Bias Contamination	1,008	2	2	0.20		
Field blank contamination	Accuracy/Bias Contamination	1,008	6	366	36.3		
Non-compliant project specific labeled analog recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	1,008	3	58	5.8		

Chlorinated Herbicides

The Phase I CSO/SWO Investigati on LSM dissolved Chlorinated Her bicide dataset is comprised of six samples with 24 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Chlorinated Herbicide analyses.

Six minor data quality issues were identified in the Phase I CS O/SWO Investigation LSM dissolved Chlorinated Herbicide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Chlorinated Herbicide LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Chlorinated Herbicide Results Affected
Non-compliant holding time	Representativeness	24	2	8	33.3
Method blank contamination	Accuracy/Bias Contamination	24	1	2	8.3
Field blank contamination	Accuracy/Bias Contamination	24	4	9	37.5
Non-compliant surrogate recovery	Overall Accuracy/Bias	24	1	2	8.3
Non-compliant column percent difference	Overall Accuracy/Bias	24	4	9	37.5
Non-compliant field duplicate relative percent difference	Precision	24	2	2	8.3

Dissolved Organic Carbon

The Phase I CSO/SWO Investigation LSM dissolved DOC dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation DOC analyses.

Two minor data quality issues were identified in the Phase I C\D/SWO Investigation LSM dissolved DOC dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
DOC LSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of DOC Results Affected
Non-compliant holding time	Representativeness	4	2	2	50.0
Field blank contamination	Accuracy/Bias	4	4	4	100
	Contamination				

Total Suspended Solids

The Phase I CSO/SWO Investigation LSM dissolved TSS dataset is comprised of six samples with six associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation TSS analyses.

Total Dissolved Solids

The Phase I CSO/SWO Investigati on LSM dissolved TDS dataset is comprised of six samples with six associated results.

No major or minor data quality issues were identified during va lidation of the CSO/SWO Investigation TDS analyses.

Low Solids Mass Particulate

Semivolatile Organic Compounds

The Phase I CSO/SWO Investigation LSM particulate SVOC dataset is comprised of four samples with 200 associated results.

One major data quality issue was identified during validation of the Phase I CSO/SWO Investigation LSM particulate SVOC analyses. The internal standards phenanthrene-d10, chrysene-d12 and/or perylene-d12 exhibited recoveries below the quality control limit. Two samples and ten results are associated with these non-compliant internal standard recoveries.

The following samples and results are associated with these non-compliant internal standard recoveries:

Sample Number	Compound Affected
	4,6-Dinitro-2-methylphenol
PR1CSOCLYLP-01B	N-nitrosodiphenylamine
	4-Bromophenyl-phenylether
	Hexachlorobenzene
	Atrazine
	Pentachlorophenol
	Carbazole
	3,3'-Dichlorobenzidine
	Di-n-octylphthalate
PR1LPDUP-01B	Di-n-octylphthalate

The identified major data quality issues are described in the table below.

Major Data Quality Issues					
Semivolatile LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected
Extremely poor internal standard recovery	Overall Accuracy/Bias	200	2	10	5.0

Five minor data quality issues were identified in the Phase I C SO/SWO Investigation LSM particulate SVOC dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Semivolatile LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected
Method blank contamination	Accuracy/Bias Contamination	200	2	4	2.0
Field blank contamination	Accuracy/Bias Contamination	200	3	5	2.5
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	200	4	7	3.5
Non-compliant project specific surrogate recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	200	4	31	15.5
Non-compliant internal standard recovery	Overall Accuracy/Bias	200	1	2	1.0

Aroclor Polychlorinated Biphenyls (PCBs)

The Phase I CSO/SWO Investigation LSM particulate Aroclor PCB d ataset is comprised of four samples with 368 associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation Aroclor PCB analyses.

Organochlorine Pesticides

The Phase I CSO/SWO Investigation LSM particulate Organochlorin e Pesticide dataset is comprised of four samples with 112 associated results.

One major data quality issue was identified during validation of the Phase I CSO/SWO Investigation LSM particulate Organochlorine Pesticide analyses. The labeled ana log 13C12-endrin aldehyde exhibited recoveries below the method quality control limit for sample PRCSOCLYLP-02B affecting the associated endrin aldehyde sample result. The identified major data quality issues are described in the table below.

Major Data Quality Issues					
Organochlorine Pesticides LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Organochlorine Pesticide Results Affected
Extremely poor method labeled	Overall	112	1	1	0.89
analog recovery	Accuracy/Bias				

Five minor data quality issues were identified in the Phase I CSO/SWO Investigation LSM particulate Organochlorine Pesticide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues	Minor Data Quality Issues					
Organochlorine Pesticides LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Organochlorine Pesticide Results Affected	
Field blank contamination	Accuracy/Bias Contamination	112	4	33	29.5	
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	112	1	3	2.7	
Non-compliant internal standard recovery	Overall Accuracy/Bias	112	4	80	71.4	
Non-compliant method labeled analog recovery	Overall Accuracy/Bias	112	1	1	0.89	
Non-compliant project specific labeled analog recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	112	4	16	14.3	

Semivolatile Organic Compounds - Select Ion Monitoring

The Phase I CSO/SWO Investigation LSM particulate SVOCs SIM dat aset is comprised of four samples with 120 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation SVOCs SIM analyses.

Seven minor data quality issues were identified in the Phase I CSO/SWO Investigation LSM particulate SVOCs SIM dataset. The identified minor data quality issues are described in the table below.

June 2016

Minor Data Quality Issues					
Semivolatiles SIM LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC SIM Results Affected
Field blank contamination	Accuracy/Bias Contamination	120	4	28	23.3
Non-compliant initial calibration relative standard deviation recovery	Overall Accuracy/Bias	120	2	2	1.7
Non-compliant project specific surrogate recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	120	1	11	9.2
Non-compliant matrix spike/matrix spike duplicate recovery	Overall Accuracy/Bias	120	1	13	10.8
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	120	1	17	14.2
Non-compliant internal standard recovery	Overall Accuracy/Bias	120	2	6	5.0
Non-compliant field duplicate relative percent difference	Precision	120	4	60	50.0

Polychlorinated Dibenzo-p-dioxins / Polychlorinated Dibenzofurans

The Phase I CSO/SWO Investigation LSM particulate PCDD/PCDFs dataset is comprised of six samples with 102 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCDD/PCDFs analyses.

Three minor data quality issues were identified in the Phase I CSO/SWO Investigation LSM particulate PCDD/PCDF dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
PCDD/PCDFs LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCDD/PCDF Results Affected
Field blank contamination	Accuracy/Bias Contamination	102	3	8	7.84
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	102	1	1	0.98
Non-compliant field duplicate relative percent difference	Precision	102	4	12	11.8

Polychlorinated Biphenyl Congeners

The Phase I CSO/SWO Investigation LSM particulate PCB Congener dataset is comprised of six samples with 1,008 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCB Congener analyses.

Six minor data quality issues were identified in the Phase I CSO/SWO Investigation LSM particulate PCB Congener dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
PCB Congeners LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCB Congener Results Affected
Method blank contamination	Accuracy/Bias Contamination	1,008	2	5	0.50
Field blank contamination	Accuracy/Bias Contamination	1,008	6	275	27.3
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	1,008	1	1	0.10
Non-compliant internal standard recovery	Overall Accuracy/Bias	1,008	4	150	14.9
Non-compliant project specific labeled analog recovery, as specified by USEPA Region 2	Overall Accuracy/Bias	1,008	3	8	0.79
Non-compliant field duplicate relative percent difference	Precision	1,008	19	19	0.88

Chlorinated Herbicides

The Phase I CSO/SWO Investigation LSM particulate Chlorinated H erbicide dataset is comprised of six samples with 24 associated results.

No major data quality issues were identified during validation of the Phase 1 CSO/SWO Investigation Chlorinated Herbicide analyses.

One minor data quality issue was identified in the Phase I CSO/ SWO Investigation LSM particulate Chlorinated Herbicide dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
Chlorinated Herbicide LSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Chlorinated Herbicide Results Affected
Non-compliant continuing	Overall	24	2	2	8.3
calibration percent difference	Accuracy/Bias				

Particulate Organic Carbon

The Phase I CSO/SWO Investigation LSM particulate POC dataset is comprised of four samples with four associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation POC analyses.

4.1.5 High Solids Mass Samples Systematic Data Quality Issues

Two systematic data quality issues were identified during the Phase I CSO/SWO Investigation High Solids Mass data validation task. These systematic data quality issues are summarized below:

- □ All field blanks contained 2,4'-DDE, 2,4'-DDD, 2,4'-DDT and 4,4'-DDT resulting in the positive results being qualified non-detected "U".
- All closing continuing calibration percent differences for Di-n-octylphthalate were outside the quality control limit. All results for Di-n-octylphthalate were qualified as estimated.

4.1.6 High Solids Mass Samples Systematic and Random Data Quality Issues by Analytical Group

High Solids Mass Dissolved

Semivolatile Organic Compounds

The Phase I CSO/SWO Investigation HSM dissolved sample SVOC dataset is comprised of four samples with 200 associated results.

One major data quality issue was identified during validation of the Phase I CSO/SWO Investigation HSM dissolved SVOC analyses. The internal standards phenanthrene-d 0 and perylene-d12 exhibited recoveries below the quality control limit. The following samples and results are associated with these non-compliant internal standard recoveries:

Sample Number	Compound Affected
	4,6-Dinitro-2-methylphenol
	N-nitrosodiphenylamine
	4-Bromophenyl-phenylether
PR1CSOCLYHD-01B	Hexachlorobenzene
PR1HDDUP-01B	Atrazine
	Pentachlorophenol
	Carbazole
	Di-n-octylphthalate

The identified major data quality issues are described in the table below.

Major Data Quality Issues					
Semivolatiles HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected
Extremely poor internal standard recovery	Overall Accuracy/Bias	200	2	16	8.0

Seven minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM dissolved SVOC dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues									
Semivolatiles HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected				
Field blank contamination	Accuracy/Bias Contamination	200	3	4	2.0				
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	200	4	14	7.0				
Non-complaint surrogate recovery	Overall Accuracy/Bias	200	4	10	5.0				
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	200	1	7	3.5				
Non-compliant internal standard recovery	Overall Accuracy/Bias	200	2	2	1.0				
Non-compliant field duplicate relative percent difference	Precision	200	2	4	2.0				
Non-compliant other quality issues	Overall Accuracy/Bias	200	1	1	0.50				

Volatile Organic Compounds (VOCs)

The Phase I CSO/SWO Investigation HSM dissolved VOC dataset is comprised of four samples with 24 associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation VOCs analyses.

Aroclor Polychlorinated Biphenyls

The Phase I CSO/SWO Investigation HSM dissolved Aroclor PCB dat aset is comprised of four samples with 36 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Aroclor PCB analyses.

One minor data quality issue was identified in the Phase I CSOSWO Investigation HSM dissolved Aroclor PCB dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
Aroclor PCBs HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Aroclor PCB Results Affected
Non-compliant surrogate recovery	Overall	36	2	18	50.0
	Accuracy/Bias				

Organochlorine Pesticides

The Phase I CSO/SWO Investigation HSM dissolved Organochlorine Pesticide dataset is comprised of four samples with 112 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Organochlorine Pesticide analyses.

Six minor data quality issues were identified in the Phase I CS O/SWO Investigation HSM dissolved Organochlorine Pesticide dataset. The identified minor data quality issues are described in the table below.

June 2016

Minor Data Quality Issues					
Organochlorine Pesticides HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Organochlorine Pesticide Results Affected
Field blank contamination	Accuracy/Bias Contamination	112	4	32	28.6
Non-compliant qualitative requirements	Overall Accuracy/Bias	112	1	1	0.89
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	112	2	6	5.4
Non-compliant internal standards	Overall Accuracy/Bias	112	4	103	92.0
Non-complaint project specific labeled analog recovery as specified by USEPA Region 2	Overall Accuracy/Bias	112	4	20	17.9
Non-compliant field duplicate relative percent difference	Precision	112	4	10	8.9

Semivolatile Organic Compounds - Select Ion Monitoring

The Phase I CSO/SWO Investigation HSM dissolved SVOCs-SIM datas et is comprised of four samples with 120 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation SVOC SIM analyses.

Six minor data quality issues were identified in the Phase I C9/SWO Investigation HSM dissolved SVOCs SIM dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues	Minor Data Quality Issues								
Semivolatiles SIM HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC SIM Results Affected				
Non-compliant holding time	Representativeness	120	2	60	50.0				
Method blank contamination	Accuracy/Bias Contamination	120	2	3	2.5				
Field blank contamination	Accuracy/Bias Contamination	120	4	35	29.2				
Non-compliant initial calibration relative standard deviation	Overall Accuracy/Bias	120	2	2	1.7				
Non-compliant project specific surrogate recovery as specified by USEPA Region 2	Overall Accuracy/Bias	120	1	16	13.3				
Non-compliant field duplicate relative percent difference	Precision	120	2	4	3.3				

Cyanide

The Phase I CSO/SWO Investigation HSM dissolved Cyanide dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Cyanide analyses.

One minor data quality issue was identified in the Phase I CSOSWO Investigation HSM dissolved Cyanide dataset. The identified minor data quality issues, and is described in the table below.

Minor Data Quality Issues					
Cyanide HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Cyanide Results Affected
Non-compliant field duplicate relative percent difference	Precision	4	2	2	50.0

Polychlorinated Dibenzo-p-dioxins / Polychlorinated Dibenzofurans

The Phase I CSO/SWO Investigation HSM dissolved PCDD/PCDFs data set is comprised of six samples with 102 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCDD/PCDFs analyses.

Five minor data quality issues were identified in the Phase I C SO/SWO Investigation HSM dissolved PCDD/PCDF dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues									
PCDD/PCDFs HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCDD/PCDF Results Affected				
Field blank contamination	Accuracy/Bias Contamination	102	2	9	8.8				
Non-compliant matrix spike/matrix spike duplicate recovery	Overall Accuracy/Bias	102	2	2	2.0				
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	102	1	1	0.98				
Non-complaint project specific labeled analog recovery as specified by USEPA Region 2	Overall Accuracy/Bias	102	6	41	40.2				
Non-compliant field duplicate relative percent difference	Precision	102	4	12	11.8				

Polychlorinated Biphenyl Congeners

The Phase I CSO/SWO Investigation HSM dissolved PCB Congener da taset is comprised of six samples with 1,008 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCB Congener analyses.

Four minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM dissolved PCB Congener dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
PCB Congeners HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCB Congener Results Affected			
Method blank contamination	Accuracy/Bias Contamination	1,008	2	2	0.20			
Field blank contamination	Accuracy/Bias Contamination	1,008	6	305	30.3			
Non-compliant internal standards	Overall Accuracy/Bias	1,008	6	400	39.7			
Non-complaint project specific labeled analog recovery as specified by USEPA Region 2	Overall Accuracy/Bias	1,008	4	72	7.1			

Chlorinated Herbicides

The Phase I CSO/SWO Investigati on HSM dissolved Chlorinated Her bicide dataset is comprised of six samples with 24 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Chlorinated Herbicides analyses.

Three minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM dissolved Chlorinated Herbicide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues									
Chlorinated Herbicide HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Chlorinated Herbicide Results Affected				
Method blank contamination	Accuracy/Bias Contamination	24	1	1	4.2				
Field blank contamination	Accuracy/Bias Contamination	24	2	7	29.2				
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	24	6	13	54.2				

Total Organic Carbon

The Phase I CSO/SWO Investigation HSM dissolved TOC dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TOC analyses.

One minor data quality issue was identified in the Phase I CSO/ SWO Investigation HSM dissolved TOC dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
TOC HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TOC Results Affected
Field blank contamination	Accuracy/Bias Contamination	4	2	2	50.0

Total Extractable Petroleum Hydrocarbons

The Phase I CSO/SWO Investigation HSM dissolved TEPH dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the CSO/SWO Investigation TEPH analyses.

Three minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM dissolved TEPH dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
TEPH HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TEPH Results Affected			
Field blank contamination	Accuracy/Bias Contamination	4	2	2	50.0			
Non-compliant initial calibration relative standard deviation	Overall Accuracy/Bias	4	2	2	50.0			
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	4	2	2	50.0			

Total Suspended Solids

The Phase I CSO/SWO Investigation HSM dissolved TSS dataset is comprised of eight samples with eight associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TSS analyses.

Two minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM dissolved TSS dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
TSS HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TSS Results Affected			
Field blank contamination	Accuracy/Bias Contamination	8	2	2	25.0			
Non-compliant field duplicate relative percent difference	Precision	8	4	4	50.0			

Total Dissolved Solids

The Phase I CSO/SWO Investigation HSM dissolved TDS dataset is comprised of eight samples with eight associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TDS analyses.

One minor data quality issue was identified in the Phase I CSO/ SWO Investigation HSM dissolved TDS dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
TDS HSM Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TDS Results Affected
Field blank contamination	Accuracy/Bias Contamination	8	2	2	25.0

High Solids Mass Particulate

Semivolatile Organic Compounds

The Phase I CSO/SWO Investigation HSM particulate SVOC dataset is comprised of four samples with 200 associated results.

One major data quality issue was identified during validation of the Phase I CSO/SWO Investigation HSM particulate SVOC analyses. The internal standard perylene-d12 exhibited recoveries below the quality control limit for samples PR1CSOCLYHP-01B and PR1HPDUP-01B associated with di-n-octylphthalate.

The identified major data quality issue is described in the table below.

Major Data Quality Issues					
Semivolatiles HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected
Extremely poor internal standard recovery	Overall Accuracy/Bias	200	2	2	1.0

Eight minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM particulate SVOC dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Semivolatiles HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC Results Affected
Non-compliant holding time	Representativeness	200	2	2	1.0
Field blank contamination	Accuracy/Bias Contamination	200	1	2	1.0
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	200	4	12	6.0
Non-compliant internal standard recovery	Overall Accuracy/Bias	200	2	2	1.0
Non-compliant method surrogate recovery	Overall Accuracy/Bias	200	3	9	4.5
Non-complaint project specific surrogate recovery as specified by USEPA Region 2	Overall Accuracy/Bias	200	4	12	6.0
Percent moisture between 50-90%	Overall Accuracy/Bias	200	4	200	100
Non-compliant linear range exceedance	Overall Accuracy/Bias	200	1	2	1.0

Volatile Organic Compounds

The Phase I CSO/SWO Investigation HSM particulate VOC dataset is comprised of seven samples with 42 associated results.

One major data quality issue was identified during validation of the Phase I CSO/SWO Investigation HSM dissolved VOC analyses. The internal standards chlorobenzene-d5 and 1,4-dichlorobenzene-d4 exhibited recoveries below the quality control limit. The following same and results are associated with these non-compliant internal standard recoveries:

Sample Number	Compound Affected
PR1CSOCLYHP-01B	1,3-Dichlorobenzene
	1,2-Dichlorobenzene
	1,2,4-Trichlorobenzene
	1,2,3-Trichlorobenzene
PR1HPDUP-01B	1,3-Dichlorobenzene
	1,2-Dichlorobenzene
	1,2,4-Trichlorobenzene
	1,2,3-Trichlorobenzene
PR1CSOCLYHP-01B-DEB	1,3-Dichlorobenzene
	1,2-Dichlorobenzene
	1,2,4-Trichlorobenzene
	1,2,3-Trichlorobenzene
PR1CSOCLYHP-02A1	1,3-Dichlorobenzene
	1,2-Dichlorobenzene
	1,2,4-Trichlorobenzene
	1,2,3-Trichlorobenzene
PR1CSOCLYHP-02A2	Chlorobenzene
	1,3-Dichlorobenzene
	1,2-Dichlorobenzene
	1,2,4-Trichlorobenzene
	1,2,3-Trichlorobenzene
PR1HPDUP-02A2	1,3-Dichlorobenzene
	1,2-Dichlorobenzene
	1,2,4-Trichlorobenzene
	1,2,3-Trichlorobenzene

The identified major data quality issue is described in the table below.

Major Data Quality Issues					
Volatiles HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of VOC Results Affected
Extremely poor internal standard recovery	Overall Accuracy/Bias	42	6	25	59.5

Six minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM particulate VOC dataset. The identified minor data quality issues are described in the table below.

Volatiles HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of VOC Results Affected
Method blank contamination	Accuracy/Bias Contamination	42	5	7	16.7
Non-compliant internal standard recovery	Overall Accuracy/Bias	42	6	10	23.8
Non-compliant surrogate recovery	Overall Accuracy/Bias	42	1	6	14.3
Non-compliant matrix spike/matrix spike duplicate recovery	Overall Accuracy/Bias	42	2	4	9.5
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	42	2	6	14.3
Percent moisture between 50-90%	Overall Accuracy/Bias	42	7	42	100

Aroclor Polychlorinated Biphenyls (PCBs)

The Phase I CSO/SWO Investigation HSM particulate Aroclor PCB dataset is comprised of four samples with 36 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Aroclor PCB analyses.

Two minor data quality issues were identified in the Phase I CS O/SWO Investigation HSM particulate Aroclor PCB dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Aroclor PCBs HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Aroclor PCB Results Affected
Non-compliant column percent	Overall	36	4	6	16.7
difference	Accuracy/Bias				
Percent moisture between 50-90%	Overall	36	4	36	100
	Accuracy/Bias				

Organochlorine Pesticides

The Phase I CSO/SWO Investigation HSM particulate Organochlorin e Pesticide dataset is comprised of four samples with 112 associated results.

One major data quality issue was identified during validation of the Phase I CSO/SWO Investigation HSM dissolved Organochlorine Pesticide analyses. The labeled analog method recoveries for 13C6-Hexachlorobenzene, 13C6-alpha-BHC, 13C6-Lindane (gamma BHC), 13C6-beta-BHC, 13C12-2,4'-DDD, 13C6-delta-BHC and/or 13C12-4,4'-DDT exhibited recoveries below the quality control limit. Two samples and six results are associated with these non-compliant labeled analog method recoveries. The following samples and results are associated with these non-compliant internal standard recoveries:

Sample Number	Compound Affected
PR1CSOCLYHP-01B	4,4'Methoxychlor
	Mirex
	Endrin Aldehyde
	Endrin Keytone
PR1HPDUP-01B	4,4'Methoxychlor
	Endrin Aldehyde

The identified major data quality issue is described in the table below.

Major Data Quality Issues					
		Total			% of
Organochlorine Pesticides	Data Quality	Number	Number	Number	Organochlorine
HSM Particulate	Parameter	of Results	of Samples	of Results	Pesticide Results
	Affected	Reported	Affected	Affected	Affected
Extremely poor labeled analog	Overall	112	2	6	5.4
method recoveries	Accuracy/Bias				

Twelve minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM particulate Organochlorine Pesticide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Organochlorine Pesticides HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Organochlorine Pesticide Results Affected
Non-compliant holding time	Representativeness	112	2	56	50.0
Field blank contamination	Accuracy/Bias Contamination	112	4	20	17.9
Method blank contamination	Accuracy/Bias Contamination	112	2	2	1.8
Non-compliant internal standard recovery	Overall Accuracy/Bias	112	4	97	86.6
Non-compliant matrix spike/matrix spike duplicate recovery	Overall Accuracy/Bias	112	1	2	1.8
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	112	1	5	4.46
Non-compliant method labeled analog recovery	Overall Accuracy/Bias	112	4	8	7.1

June 2016

Non-complaint project	Overall	112	4	44	39.3
specific labeled analog	Accuracy/Bias				
recovery as specified by					
USEPA Region 2					
Non-compliant qualitative	Overall	112	2	2	1.8
requirements	Accuracy/Bias				
Non-compliant linear range	Overall	112	2	4	3.6
exceedance	Accuracy/Bias				
Percent moisture between 50-	Overall	112	4	112	100
90%	Accuracy/Bias				
Non-compliant field duplicate	Precision	112	4	34	30.4
relative percent difference					

Semivolatile Organic Compounds - Select Ion Monitoring

The Phase I CSO/SWO Investigation HSM particulate SVOCs SIM dat aset is comprised of four samples with 120 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation SVOCs SIM analyses.

Five minor data quality issues were identified in the Phase I C SO/SWO Investigation HSM particulate SVOCs SIM dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues					
Semivolatiles SIM HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of SVOC SIM Results Affected
Non-compliant holding time	Representativeness	120	2	60	50.0
Field blank contamination	Accuracy/Bias Contamination	120	2	8	6.7
Non-compliant initial calibration relative standard deviation	Overall Accuracy/Bias	120	2	2	1.7
Percent moisture between 50-90%	Overall Accuracy/Bias	120	4	120	100
Non-compliant field duplicate relative percent difference	Precision	120	4	12	10.0

Cyanide

The Phase I CSO/SWO Investigation HSM particulate Cyanide dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Cyanide analyses.

Three minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM particulate Cyanide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues						
Cyanide HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Cyanide Results Affected	
Field blank contamination	Accuracy/Bias Contamination	4	3	3	75.0	
Non-compliant matrix spike/matrix spike duplicate recovery	Overall Accuracy/Bias	4	1	1	25.0	
Percent moisture between 50-90%	Overall Accuracy/Bias	4	4	4	100	

Polychlorinated Dibenzo-p-dioxins / Polychlorinated Dibenzofurans

The Phase I CSO/SWO Investigation HSM particulate PCDD/PCDFs dataset is comprised of six samples with 102 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCDD/PCDFs analyses.

Four minor data quality issues were identified in the Phase I C SO/SWO Investigation HSM particulate PCDD/PCDFs dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
PCDD/PCDFs HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCDD/PCDF Results Affected		
Field blank contamination	Accuracy/Bias Contamination	102	3	5	4.9		
Non-compliant internal standard recovery	Overall Accuracy/Bias	102	2	20	19.6		
Percent moisture between 50-90%	Overall Accuracy/Bias	102	4	68	66.7		
Non-compliant field duplicate relative percent difference	Precision	102	2	4	3.9		

Polychlorinated Biphenyl Congeners

The Phase I CSO/SWO Investigation HSM particulate PCB Congener dataset is comprised of six samples with 1,008 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation PCB Congeners analyses.

Eight minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM particulate PCB Congener dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues						
PCB Congeners HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of PCB Congener Results Affected	
Method blank contamination	Accuracy/Bias Contamination	1,008	4	10	0.99	
Field blank contamination	Accuracy/Bias Contamination	1,008	3	22	2.2	
Non-compliant matrix spike/matrix spike duplicate relative percent difference	Precision	1,008	1	1	0.10	
Non-compliant internal standard recovery	Overall Accuracy/Bias	1,008	5	413	41.0	
Non-compliant method labeled analog recovery	Overall Accuracy/Bias	1,008	1	1	0.10	
Non-complaint project specific labeled analog recovery as specified by USEPA Region 2	Overall Accuracy/Bias	1,008	5	49	4.9	
Percent moisture between 50-90%	Overall Accuracy/Bias	1,008	4	672	66.7	
Non-compliant field duplicate relative percent difference	Precision	1,008	4	40	4.0	

Chlorinated Herbicides

The Phase I CSO/SWO Investigation HSM particulate Chlorinated H erbicide dataset is comprised of six samples with 24 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Chlorinated Herbicides analyses.

Eight minor data quality issues were identified in the Phase I CSO/SWO Investigation HSM particulate Chlorinated Herbicide dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues							
Chlorinated Herbicide HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Chlorinated Herbicide Results Affected		
Method blank contamination	Accuracy/Bias Contamination	24	3	4	16.7		
Field blank contamination	Accuracy/Bias Contamination	24	6	10	42.0		

June 2016

Non-compliant matrix spike/matrix	Overall	24	3	11	45.8
spike duplicate recovery	Accuracy/Bias				
Non-compliant matrix spike/matrix	Precision	24	3	7	29.2
spike duplicate relative percent					
difference					
Non-compliant surrogate recovery	Overall	24	1	4	16.7
	Accuracy/Bias				
Non-compliant laboratory control	Overall	24	2	4	16.7
standard recovery	Accuracy/Bias				
Non-compliant column percent	Overall	24	4	10	41.7
difference	Accuracy/Bias				
Percent moisture between 50-90%	Overall	24	6	24	100
	Accuracy/Bias				

Total Organic Carbon

The Phase I CSO/SWO Investigation HSM particulate TOC dataset i s comprised of six samples with six associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TOC analyses.

One minor data quality issue was identified in the Phase I CSO/SWO Investigation HSM particulate TOC dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
TOC HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TOC Results Affected
Percent moisture between 50-90%	Overall Accuracy/Bias	6	6	6	100

Total Petroleum Hydrocarbons

The Phase I CSO/SWO Investigation HSM particulate TEPH dataset is comprised of four samples with four associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TEPH analyses.

Five minor data quality issue was identified in the Phase I CSOSWO Investigation HSM particulate TEPH dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
TEPH HSM Particulate	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TEPH Results Affected			
Non-compliant holding time	Representativeness	4	2	2	50.0			
Non-compliant initial calibration relative standard deviation	Overall Accuracy/Bias	4	2	2	50.0			
Non-compliant continuing calibration percent difference	Overall Accuracy/Bias	4	2	2	50.0			
Percent moisture between 50-90%	Overall Accuracy/Bias	4	4	4	100			
Non-compliant field duplicate relative percent difference	Precision	4	2	2	50.0			

4.1.7 Grab Water Samples Systematic Data Quality Issues

No systematic data quality issues were identified during the Ph ase I CSO/SWO Investigation data grab water sample validation task.

4.1.8 Grab Water Samples Systematic and Random Data Quality Issues by Analytical Group

Grab Water

Metals

The Phase I CSO/SWO Investigation grab water sample Metals dataset is comprised of four samples with 92 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Metals analyses.

Two minor data quality issues were identified in the Phase I CS O/SWO Investigation grab water Metals dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues									
Metals Grab Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Metals Results Affected				
Continuing calibration blank contamination	Accuracy/Bias Contamination	92	2	6	6.5				
Non-compliant field duplicate relative percent difference	Precision	92	2	8	8.7				

Mercury

The Phase I CSO/SWO Investigation grab water sample Mercury dataset is comprised of four samples with four associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation Mercury analyses.

Methyl Mercury

The Phase I CSO/SWO Investigation grab water sample Methyl Merc ury dataset is comprised of four samples with four associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation Methyl Mercury analyses.

Total Suspended Solids

The Phase I CSO/SWO Investigation grab water sample TSS dataset is comprised of 45 samples with 45 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TSS analyses.

One minor data quality issue was identified in the Phase I CSOSWO Investigation grab water TSS dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues								
TSS Grab Water	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of TSS Results Affected			
Non-compliant holding time	Representativeness	45	8	8	17.8			

Total Dissolved Solids

The Phase I CSO/SWO Investigation grab water sample TDS dataset is comprised of 45 samples with 45 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation TDS analyses.

One minor data quality issue was identified in the Phase I CSOSWO Investigation grab water TDS dataset. The identified minor data quality issue is described in the table below.

Minor Data Quality Issues					
TDS		Total			% of
Grab Water	Data Quality	Number	Number	Number	TDS
	Parameter	of Results	of Samples	of Results	Results
	Affected	Reported	Affected	Affected	Affected
Non-compliant holding time	Representativeness	45	8	8	17.8

Grab Water Dissolved

Metals

The Phase I CSO/SWO Investigation grab water dissolved sample M etals dataset is comprised of four samples with 92 associated results.

No major data quality issues were identified during validation of the Phase I CSO/SWO Investigation Metals analyses.

Three minor data quality issues were identified in the Phase CSO/SWO Investigation dissolved grab water Metals dataset. The identified minor data quality issues are described in the table below.

Minor Data Quality Issues								
Metals Grab Water Dissolved	Data Quality Parameter Affected	Total Number of Results Reported	Number of Samples Affected	Number of Results Affected	% of Metals Results Affected			
Field blank contamination	Accuracy/Bias Contamination	92	4	8	8.7			
Continuing calibration blank contamination	Accuracy/Bias Contamination	92	4	9	9.8			
Non-compliant field duplicate relative percent difference	Precision	92	2	2	2.2			

Mercury

The Phase I CSO/SWO Investigation grab water dissolved sample M ercury dataset is comprised of four samples with four associated results.

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation Mercury analyses.

Methyl Mercury

The Phase I CSO/SWO Investigation grab water dissolved sample Methyl Mercury dataset is comprised of four samples with four associated results.

CSO/SWO Phase I Data Quality Usability Assessment Report – Rev 2

June 2016

No major or minor data quality issues were identified during va lidation of the Phase I CSO/SWO Investigation Methyl Mercury analyses.

5. Total Tetrachlorinated Dibenzo-p-dioxin Verification

This verification procedure was implemented as an evaluation of Total Tetrachlorinated Dibenzo-p-Dioxin (TCDD) results since these values were not evaluated during the isomer specific data validation task. This process is used to assess both the completeness and accuracy of the total TCDD data set.

Total TCDD results were verified for each sample having total T CDD results reported in Phase I of the CSO/SWO Investigation. In cases where multiple analyses were performed by the laboratory for 2,3,7,8-TCDD (example: multiple dilutions due to elevated target analyt e concentrations or re-analysis based on failed quality control criteria), EDS staff made certain that the total TCDD value reported in the data base, as well as hardcopy data, was based on the same analysis used to derive the 2,3,7,8-TCDD value reported.

Procedure Acceptance Criteria:

	Selected ion current profiles (SICPs) for ions 319.8965 and 321.8936 representing all non 2,3,7,8-substituted tetra chlorinated dibenzo-p-dioxins and 2,3,7,8-substituted tetra chlorinated dibenzo-p-dioxin are reported for each sample.
	Integrated areas are present for both the primary and confirmation ions for all peaks and are 2.5 times above background noise in each sample SICP.
	Instrument quantitation reports containing relative response factors for 2,3,7,8-TCDD, area counts for the 2,3,7,8 –TCDD labeled analog and sample preparation information are presen for each sample.
Cal	culation Acceptance Criteria:
	The retention time of each non 2,3,7,8-substituted compound identified as present in the sample was within the window established by the window defining mixture, for the tetra chlorinated homologue.
	The integrated ion current of each non 2,3,7,8-substituted compound identified as present in the sample was at least 2.5 times background noise.
	All peaks meeting the requirements described above were included in the laboratory's calculation of Total TCDD.

Results of Verification:

concentration recalculated.

All 53 total TCDD results, reported during implementation of the Phase I CSO/SWO Investigation, were evaluated during this task. Of the 53 samples evaluated for this program, four of the results are recommended for editing based on the results of the total TCDD result verification task. The affected samples and associated results are provided in Table 5-1 below. Total TCDD results for

A minimum of one non 2,3,7,8-substituted compound identified was verified and the

2,3,7,8-substituted tetra chlorinated dibenzo-p-dioxin identified in each sample.

Recalculate the sum of all non 2,3,7,8-substituted tetra chlorinated dibenzo-p-dioxins and

these samples have been corrected in both the laboratory hardcopy data reports and United States Environmental Protection Agency (USEPA) Region 2 Main Electronic Data Deliverable (MEDD).

Table 5-1

Sample Identification	Result Units	Existing Result Value	Data Qualifiers	New Result Value	Data Oualifiers
PR1LPDUP-01A	pg/g	11.5	EMPC	9.72	EMPC
PR1CSOCLYHP-02B PR1HPDUP-02B	pg/g pg/g	14.0	EMPC	12.8	EMPC
PR1CSOCLYHP-01C	pg/g	19.4	EMPC	17.8	EMPC

6. Conclusions

The data usability evaluations outlined in this report provides details regarding the relationship of data quality issues to associated samples and sample results. Ninet y-nine percent of the data validated and reported are suitable for their intended use. A total of 29 sa mple results for the SVOC analyses and 25 sample results for the VOC analyses were rejected due to intern all standard recoveries. A total of seven sample results for the organochlorine pesticide analyses were rejected due to method labeled analog recoveries. Sample results that were rejected are not suitable for project use. Sample results that are qualified as estimated due to multiple minor data quality issues as detailed in this report are suitable for project use. The achievement of the completeness goals for number of samples collected and the number of samples accepted for use provides sufficient quality data to support project decisions.

7. References

Tierra 2013. Combined Sewer Overflow/Stormwater Outfall Investigation Quality Assurance Project Plan, Revision 3, September.

Attachment 1 Phase I Report Addendum – Additional Data Evaluation

Combined Sewer Overflow/Stormwater Outfall Investigation

Phase I Report Addendum – Additional Data Evaluation

Tierra Solutions, Inc.

East Brunswick, New Jersey

March 2016

Revision 0

1.	Intro	duction	1-1
	1.1	Additional Data Evaluation Process	1-2
		1.1.1 Unit Conversion	1-3
		1.1.2 Chi-Square Test	1-4
2.	Addit	ional Data Evaluation Findings	2-1
	2.1	Polychlorinated Dibenzo-p-Dioxins/Polychlorinated Dibenzofurans	2-2
	2.2	Polychlorinated Biphenyl Congeners	2-3
	2.3	Polychlorinated Biphenyl Aroclors	2-4
	2.4	Organochlorine Pesticide	2-4
	2.5	Semivolatile Organic Compounds	2-5
	2.6	Semivolatile Organic Compounds Selective Ion Monitoring	2-6
	2.7	Chlorinated Herbicides	2-7
	2.8	Cyanide	2-8
	2.9	Volatile Organic Compounds	2-9
	2.10	Total Extractable Petroleum Hydrocarbons	2-9
3.	Sumr	mary	3-1
4.	Refer	ences	4-1
Арре	ndices	5	
Α		Data Evaluation Summaries and Analytical Results – PCDDs/PCDFs	
В		Data Evaluation Summaries and Analytical Results – PCB Congeners	
C Data Evaluation Summaries a		Data Evaluation Summaries and Analytical Results – Aroclor PCBs	
D		Data Evaluation Summaries and Analytical Results – Organochlorine Pesticide	
Е		Data Evaluation Summaries and Analytical Results – SVOCs	
F		Data Evaluation Summaries and Analytical Results – SVOC SIM	
G		Data Evaluation Summaries and Analytical Results – Chlorinated Herbicides	

i

Table of Contents

Н	Data Evaluation Summaries and Analytical Results – Cyanide
l	Data Evaluation Summaries and Analytical Results – VOCs
.1	Data Evaluation Summaries and Analytical Results – TEPH

1. Introduction

This Phase I Report Addendum – Additional Data Evaluation (Phase I Report Addendum) has been developed by Tierra Solutions, Inc. (Tierra), on behalf of Occidental Chemical Corporation, the successor to Diamond Shamrock Chemicals Company (formerly known as Diamond Alkali Company). Tierra prepared the Phase I Evaluation/Recommendation Report (Phase I Report, Revision 0; Tierra 2014) to document the data evaluation completed as part of Phase I of the combined sewer overflow/stormwater outfall (CSO/SWO) investigation implemented under the U.S. Environmental Protection Agency- (USEPA-) approved Combined Sewer Overflow/Stormwater Outfall Investigation Quality Assurance Project Plan (CSO/SWO Investigation QAPP; Tierra 2013). In response to USEPA comments (specifically, Comment No. 3), dated August 6, 2015 on the Phase I Report (Revision 0; Tierra 2014), Tierra conducted additional evaluations of the Phase I CSO sampling results/data. These additional data evaluations were beyond the scope of the data evaluation criteria defined in the CSO/SWO Investigation QAPP (Tierra 2013).

The Phase I data evaluation was conducted on an analytical group basis, for each sampling method, and was designed to identify the most sensitive sampling method by comparing the number of detections of target analytes within a given analytical group. However, in order to address USEPA comment No. 3, Tierra conducted additional data evaluations by tabulating the results from the high solids mass (HSM), low solids mass (LSM), and whole water datasets in terms of both concentration and frequency of detections, developing summary statistics, and reviewing the results for trends to determine if any new insights could be gathered to help in the planning for Phase II of the CSO/SWO program. The additional data evaluations consisted of side-by-side comparisons of the HSM and LSM particulate phases, dissolved-phases, total concentrations, and whole water total concentrations detected in the samples collected during Phase I. This Phase I Report Addendum documents the additional data evaluation methods, summary statistics, and results associated with Phase I of the CSO/SWO investigation.

Preliminary results of the additional data evaluations and summary statistics (for select analytical groups) were presented to the USEPA in a meeting on November 17, 2015, and this Phase I Report Addendum provides the results of the additional data evaluations as requested by the USEPA. Additional data evaluations were completed for the following analytical groups:

- AMERICA	Polychlorinated dibenzo-p-dioxins/polychlorinated dibenzofurans (PCDDs/PCDFs)
-	Polychlorinated biphenyl (PCB) congeners
and the same of th	PCB Aroclors
-	Organochlorine pesticides (OCPs)

	Semivolatile organic compounds (SVOCs)							
Ш	SVOC selective ion monitoring (SIM)							
Ш	Chlorinated herbicides							
Ш	Cyanide (CN)							
Ш	Volatile organic compounds (VOCs)							
Ш	Total extractable petroleum hydrocarbons (TEPH)							
1.1	Additional Data Evaluation Process							
fred	e additional data evaluations performed by Tierra included detailed statistical analyses to compare the quency of detections and differences in concentrations between each sampling method used in Phase I SM, LSM, and whole water). Specific details of the evaluations performed include the following:							
	Comparison of the frequency of detections Between the whole water sampling method and HSM total and LSM total sampling methods. HSM total and LSM total were estimated as the sum of the HSM or LSM particulate concentration (e.g., micrograms per kilogram [µg/kg]) and the corresponding dissolved-phase concentration (e.g., micrograms per liter [µg/L]). Additional details on HSM and LSM total calculation and unit conversion are provided in Section 1.1.1. Between the HSM and LSM sampling methods for both the particulate and dissolved-phases.							
	 Comparison of constituent concentrations Between sampling methods for analytes detected by two or more sampling methods for whole water. Between HSM (total, particulate, and dissolved-phase) and LSM (total, particulate, and dissolved-phase). 							
The	e following data evaluation rules were applied for each analytical group:							
	Only positively identified analytes (results reported above the project quantitation limit [PQL]) were included in the additional data evaluation.							
Ш	Analytes reported as non-detects were assigned a zero value for both the statistical analyses, as well as for averaging primary and duplicate sample concentrations.							

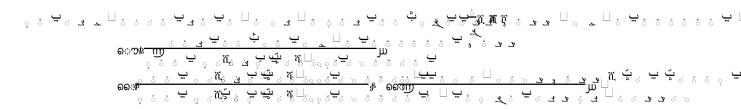
Analytes identified above the method detection limit but below the PQL were assigned a zero value for both the statistical analyses, as well as for averaging primary and duplicate sample concentrations.

1.1.1 Unit Conversion

To perform a side-by-side comparison of the HSM and LSM particulate and dissolved-phase concentrations and whole water concentrations, the particulate results reported for the HSM particulate and LSM particulate sampling methods (e.g., µg/kg) were converted to a volumetric concentration (e.g., µg/L). Converting all sample results into consistent units (e.g., mass per volume units [µg/L]) allows direct comparison of sample concentrations between sampling methods for analytes detected by two or more sampling methods.

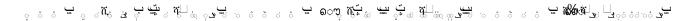
The following equations were used to convert HSM and LSM particulate results to volumetric concentrations:

HSM Particulate:



where:

- Particulate Contaminant Mass is the HSM particulate sample result (e.g., μg/kg) reported by the analytical laboratories on a mass per unit weight (dry weight) basis
- ☐ Total Solids Mass (wet weight) refers to the total solids sample mass collected for each event as presented in Table 2-1
- ☐ Total Liters Processed refers to the total liters of CSO overflow processed for each event as presented in Table 2-1
- Unit weight (dry and wet weight) of sediment are sample-specific weights reported by the analytical laboratories and



Wet weight and percent solids information was obtained for each analytical group for each event/attempt from the analytical laboratories.

LSM Particulate:



where:

- Particulate Contaminant Mass is the HSM particulate sample result (e.g., μg/kg) reported by the analytical laboratories on a mass per unit weight (dry weight) basis

 Solids Mass on Filter (dry weight) refers to the sample-specific solids mass collected on the filter during LSM filtration and is based on total suspended solids and total LSM bulk sample volume filtered

 Total LSM Bulk Volume Filtered refers to the total LSM bulk sample volume filtered (sample-specific) to
- ☐ Total LSM Bulk Volume Filtered refers to the total LSM bulk sample volume filtered (sample-specific) to generate LSM particulate (on the filter) and LSM dissolved (filtrate) samples for analysis.

Any factors needed for unit conversion were added to equations as appropriate. Additionally, the converted HSM and LSM particulate concentrations were summed with the corresponding dissolved-phase concentrations to calculate HSM total and LSM total concentrations.

1.1.2 Chi-Square Test

As an additional data evaluation step, a quantitative evaluation (statistical comparison) of the number of detected compounds in total concentrations (HSM total, LSM total, and whole water) among sampling methods was conducted. This is an additional line of comparison to evaluate if the number of detections are significantly different between sampling methods. A statistical test was applied in a pairwise manner for each sampling method and event/attempt to evaluate if the number of compounds detected within an analytical group was dependent on the sampling method. The number of detects and non-detects for each sampling method within the analytical group were arranged in a two-way contingency table (Agresti 1990). The "null hypothesis" of the test is that the frequency of detects is independent of the sampling method. When the p-value of the test is less than 0.05, the null hypothesis is rejected and the assumption is that the frequency of detection is dependent upon the sampling method (i.e., indicating that the number of detects is significantly different between methods). When the frequency within all cells of the two-way contingency table exceeded 5, a Pearson chi-squared test of independence was conducted (Agresti 1990). When the frequency in any of the cells of the two-way contingency table was less than 5, a Fisher's Exact Test (Agresti 1990) was used to test independence. Results of the chi-square test for each analytical group are presented in Section 2.

2. Additional Data Evaluation Findings

A summary of sampling events/attempts and the analytical groups selected for additional data evaluations is summarized in Table 2-1 (below).

Table 2-1
Summary of Samples Collected and Analyzed and the Volumes and Mass Associated with HSM Sampling

Event and Attempt	Sample Identification	Date	Total HSM Particulate Mass Collected (grams wet weight)	Total CSO Volume Processed (liters)	Analytical Group Selected for Additional Data Evaluation
Event 1, Attempt 1	PR1CSOCLY**-01A PR1CSOCLY**DUP- 01A	6/10/2013	223.35	13,058	PCDDs/PCDFs, PCB congeners
Event 1, Attempt 2	PR1CSOCLY**-01B PR1CSOCLY**- DUP-01B	7/1/2013	1,564	17,589	PCB Aroclors, OCPs, SVOCs, SVOC SIM, chlorinated herbicides, CN, VOCs, TEPH
Event 1, Attempt 3	PR1CSOCLY**-01C PR1CSOCLY**- DUP-01C	4/30/2014	1,575.73	14,307	PCDDs/PCDFs, PCB congeners, chlorinated herbicides,
Event 2, Attempt 1	PR1CSOCLY**-02A PR1CSOCLY**- DUP-02A	10/7/2013	219.78	1,457	VOCs
Event 2, Attempt 2	PR1CSOCLY**-02B PR1CSOCLY**- DUP-02B	12/7/2013	1,185.05	13,353	PCDDs/PCDFs, PCB congeners, PCB Aroclors, OCPs, SVOCs, SVOC SIM, chlorinated herbicides, CN, TEPH

Notes:

HSM particulate solids mass represents the total solids mass generated within the continuous flow centrifuge (CFC) during each sampling event/attempt.

Total volume of CSO processed is the total CSO volume pumped and processed through the CSO sampling system, including the CFC, LSM, and whole water sample ports during each sampling event/attempt.

^{** =} Two-character code to indicate sample matrix (e.g., "HP" for HSM particulate, "WW" for whole water).

Results of the additional data evaluations for each analytical group is summarized below. Supporting information is presented in Appendices A through J.

All three sample collection and processing methods (HSM, LSM, and whole water) were evaluated for the

2.1 Polychlorinated Dibenzo-p-Dioxins/Polychlorinated Dibenzofurans

PCDD/PCDF analytical group. Samples (primary sample and field duplicate sample) were collected for PCDD/PCDF analysis during three events: Event #1/Attempt #1, Event #1/Attempt #3, and Event #2/Attempt #2. A summary of the findings of the additional data evaluations for PCDD/PCDF data are provided below. Data evaluation summaries and analytical results are presented in Appendix A. The HSM sampling method resulted in a higher frequency of detects (number of detections) than other methods. The average frequency of detected congeners for the HSM sampling method over all events was: total – 77% (13 detects out 17 congeners), particulate – 77% (13 detects out of 17 congeners), and dissolved – 25% (four detects out of 17 congeners) (Table A-1). Where detected in both the HSM and LSM methods, the HSM total concentrations were higher (23%), on average, than the LSM total concentrations (Table A-1). Where detected in both the HSM and whole water methods, the HSM total concentrations were slightly lower (-10%), on average, than the whole water concentrations; however, there was great variability among events (Table A-1). □ Where detected in both the LSM and whole water methods, the LSM total concentrations were lower (-18%), on average, than the whole water concentrations (Table A-1). Where detected in both the HSM and LSM particulate sampling methods, the HSM particulate concentrations were lower (-48%), on average, than the LSM particulate concentrations (Table A-1). Where detected in both the HSM and LSM dissolved sampling methods, the HSM dissolved concentrations were significantly higher (501%), on average, than the LSM dissolved concentrations (Table A-1). Results of the chi-square test indicated the following:

The HSM sampling method had a significantly greater frequency of detected congeners than both of the

other methods (i.e., LSM and whole water) for all events (Table A-2).

Ш	The LSM and whole water sampling methods were similar with respect to the number of detected congeners (Table A-2).
Tab	oles A-3 through A-7 provide the analytical results and conversions for each method and sampling event.
2.2	Polychlorinated Biphenyl Congeners
PCI PCI #2/	three sample collection and processing methods (HSM, LSM, and whole water) were evaluated for the B congener analytical group. Samples (primary sample and field duplicate sample) were collected for B congener analysis during three events: Event #1/Attempt #1, Event #1/Attempt #3, and Event Attempt #2. A summary of the findings of the additional data evaluations for PCB congener data are vided below. Data evaluation summaries and analytical results are presented in Appendix B.
	The HSM sampling method resulted in a higher frequency of detects (number of detections) than other methods. The average frequency of detected congeners for the HSM sampling method over all events was: total – 59% (99 detects out 168 congeners/coelutions), particulate – 59% (99 detects out of 168 congeners/coelutions), and dissolved – 15% (25 detects out of 168 congeners/coelutions) (Table B-1).
	Where detected in both the HSM and LSM methods, the HSM total concentrations were higher (19%), on average, than the LSM total concentrations (Table B-1).
	Where detected in both the HSM and whole water methods, the HSM total concentrations were slightly lower (-10%), on average, than the whole water concentrations; however, there is great variability among events (Table B-1).
	Where detected in both the LSM and whole water methods, the LSM total concentrations were lower (-33%), on average, than the whole water concentrations (Table B-1).
	Where detected in both the HSM and LSM particulate methods, the HSM particulate concentrations were slightly lower (-2%), on average, than the LSM particulate concentrations (Table B-1).
Ш	Where detected in both the HSM and LSM dissolved methods, the HSM dissolved concentrations were higher (71%), on average, than the LSM dissolved concentrations (Table B-1).
Res	sults of the chi-square test indicated the following:
	The HSM sampling method had a higher frequency of detected congeners than both of the other methods (i.e., LSM and whole water) for all events; however, the difference for Event #1/Attempt #1 is not significant (i.e., p>0.05), with respect to the whole water method (Table B-2).

Ш	The whole water sampling method had a higher frequency of detected congeners than the LSM method for all events (Table B-2).
Tab	oles B-3 through B-7 provide the analytical results and conversions for each method and sampling event.
2.3	Polychlorinated Biphenyl Aroclors
PC Aro find	three sample collection and processing methods (HSM, LSM, and whole water) were evaluated for the B Aroclor analytical group. Samples (primary sample and field duplicate sample) were collected for PCB clor analysis during two events: Event #1/Attempt #2 and Event #2/Attempt #2. A summary of the lings of the additional data evaluations for PCB Aroclor data are provided below. Data evaluation numeries and analytical results are presented in Appendix C.
	Two PCB Aroclors (Aroclor 1254 and Aroclor 1260) were identified for HSM particulate analysis; however, only Aroclor 1254 was detected above the PQL during analysis.
	Concentration comparisons were not performed between sampling methods because PCB Aroclors were positively identified (above the PQL) for only HSM particulate analysis.
Res	sults of the chi-square test indicated the following:
Ш	There was no significant difference in frequency of detection among methods (HSM, LSM, and whole water) according to the Fisher Exact Test (Table C-2).
Tab	oles C-3 through C-7 provide the analytical results and conversions for each method and sampling event.
2.4	Organochlorine Pesticide
OC ana ado	three sample collection and processing methods (HSM, LSM, and whole water) were evaluated for the P analytical group. Samples (primary sample and field duplicate sample) were collected for OCP alysis during two events: Event #1/Attempt #2 and Event #2/Attempt #2. A summary of the findings of the litional data evaluations for OCP data are provided below. Data evaluation summaries and analytical ults are presented in Appendix D.
	The HSM sampling method resulted in a higher frequency of detects (number of detections) than other methods. The average frequency of detected congeners for the HSM sampling method over all events was: total – 45% (13 detects out 28 congeners), particulate – 35% (9.8 detects out of 28 congeners), and dissolved – 35% (9.8 detects out of 28 congeners) (Table D-1).

Ш	Where detected in both the HSM and LSM methods, the HSM total concentrations were higher (8%), or average, than the LSM total concentrations (Table D-1).
and a second	Where detected in both the HSM and whole water methods, the HSM total concentrations were slightly lower (-5%), on average, than the whole water concentrations; however, there was great variability among events (Table D-1).
	Where detected in both the LSM and whole water methods, the LSM total concentrations were slightly lower (-7%), on average, than the whole water concentrations (Table D-1).
Ш	Where detected in both the HSM and LSM particulate sampling methods, the HSM particulate concentrations were lower (-55%), on average, than the LSM particulate concentrations (Table D-1).
Ш	Where detected in both the HSM and LSM dissolved sampling methods, the HSM dissolved concentrations were higher (91%), on average, than the LSM dissolved concentrations (Table D-1).
Re	sults of the chi-square test indicated the following:
Ш	There was no significant difference in frequency of detection among methods (HSM, LSM, and whole water) (Table D-2).
Tal	oles D-3 through D-7 provide the analytical results and conversions for each method and sampling event.
2.5	Semivolatile Organic Compounds
SV ana ado	three sample collection and processing methods (HSM, LSM, and whole water) were evaluated for the OC analytical group. Samples (primary sample and field duplicate sample) were collected for SVOC alysis during two events: Event #1/Attempt #2 and Event #2/Attempt #2. A summary of the findings of the ditional data evaluations for SVOC data are provided below. Data evaluation summaries and analytical ults are presented in Appendix E.
- AMP (A)	The HSM sampling method resulted in a higher frequency of detects (number of detections) than other methods. The average frequency of detected congeners for the HSM sampling method over all events was: $total - 9\%$ (4.3 detects out of 50 compounds), particulate $- 5\%$ (2.5 detects out of 50 compounds), and dissolved $- 6\%$ (2.8 detects out of 50 compounds) (Table E-1).
Ш	Where detected in both the HSM and LSM methods, the HSM total concentrations were higher (51%), on average, than the LSM total concentrations (Table E-1).

	Where detected in both the HSM and whole water methods, the HSM total concentrations were higher (19%), on average, than the whole water concentrations (Table E-1).	
Ш	Where detected in both the LSM and whole water methods, the LSM total concentrations were lower (-33%), on average, than the whole water concentrations (Table E-1).	
	Where detected in both the HSM and LSM particulate sampling methods, the HSM particulate concentrations were lower (-81%), on average, than the LSM particulate concentrations (Table E-1).	
	Where detected in both the HSM and LSM dissolved methods, the HSM dissolved concentrations were higher (37%), on average, than the LSM dissolved concentrations (Table E-1).	
Res	sults of the chi-square test indicated the following:	
	There was some evidence that the HSM sampling method had a greater frequency of detected SVOCs than both of the other methods (i.e., LSM and whole water) for all events; however, this apparent difference was not statistically significant (i.e., p>0.05) (Table E-2).	
Ш	There was no significant difference in frequency of detection between the LSM and whole water methods (Table E-2).	
Tab	oles E-3 through E-7 provide the analytical results and conversions for each method and sampling event.	
2.6	Semivolatile Organic Compounds Selective Ion Monitoring	
SV SIM the	three sample collection and processing methods (HSM, LSM, and whole water) were evaluated for the OC SIM analytical group. Samples (primary sample and field duplicate sample) were collected for SVOC If analysis during two events: Event #1/Attempt #2 and Event #2/Attempt #2. A summary of the findings of additional data evaluations for SVOC SIM data are provided below. Data evaluation summaries and allytical results are presented in Appendix F.	
	ere detected in both the LSM and whole water methods, the LSM total concentrations were lower 3%), on average, than the whole water concentrations (Table E-1). Here detected in both the HSM and LSM particulate sampling methods, the HSM particulate ecentrations were lower (-81%), on average, than the LSM particulate concentrations (Table E-1). Here detected in both the HSM and LSM dissolved methods, the HSM dissolved concentrations were here (37%), on average, than the LSM dissolved concentrations (Table E-1). Here detected in both the HSM and LSM dissolved methods, the HSM dissolved concentrations were here (37%), on average, than the LSM dissolved concentrations (Table E-1). Here detected in both the HSM amd LSM dissolved methods, the HSM dissolved concentrations were here (37%), on average, than the LSM dissolved concentrations (Table E-1). Here was some evidence that the HSM sampling method had a greater frequency of detected SVOCs in both of the other methods (i.e., LSM and whole water) for all events; however, this apparent erence was not statistically significant (i.e., p>0.05) (Table E-2). Here was no significant difference in frequency of detection between the LSM and whole water thods (Table E-2). Here was no significant difference in frequency of detection between the LSM and whole water thods (Table E-2). Here was no significant difference in frequency of detection between the LSM and whole water thods (Table E-2). Here was no significant difference in frequency of detections for each method and sampling event. Here was no significant difference in frequency of detections for each method and sampling event. Here was no significant difference in frequency of detected sample) were evaluated for the silk analytical group. Samples (primary sample and field duplicate sample) were collected for SVOC alysis during two events: Event #1/Attempt #2 and Event #2/Attempt #2. A summary of the findings of litional data evaluations for SVOC SIM data are provided below. Data evaluation summaries and resul	
	Where detected in both the HSM and LSM methods, the HSM total concentrations were lower (-37%), on average, than the LSM total concentrations (Table F-1)	

	Where detected in both the HSM and whole water methods, the HSM total concentrations were lower (-60%), on average, than the whole water concentrations (Table F-1).
- AMERICAN	Where detected in both the LSM and whole water methods, the LSM total concentrations were lower (-27%), on average, than the whole water concentrations (Table F-1).
	Where detected in both the HSM and LSM particulate sampling methods, the HSM particulate concentrations were lower (-83%), on average, than the LSM particulate concentrations (Table F-1).
diament of the state of the sta	Where detected in both the HSM and LSM dissolved sampling methods, the HSM dissolved concentrations were higher (92%), on average, than the LSM dissolved concentrations (Table F-1).
Re	sults of the chi-square test indicated the following:
	The HSM sampling method had a greater frequency of detected SVOC SIM than both of the other methods (i.e., LSM or whole water) for all events (Table F-2).
	There was no significant difference in the frequency of detection between the LSM and whole water methods (Table F-2).
Tal	oles F-3 through F-7 provide the analytical results and conversions for each method and sampling event.
2.7	Chlorinated Herbicides
chle for Eve	three sample collection and processing methods (HSM, LSM, and whole water) were evaluated for the orinated herbicides analytical group. Samples (primary sample and field duplicate sample) were collected chlorinated herbicides analysis during three events: Event #1/Attempt #2, Event #1/Attempt #3, and ent #2/Attempt #2. A summary of the findings of the additional data evaluations for chlorinated herbicides a are provided below. Data evaluation summaries and analytical results are presented in Appendix G.
AND CO.	The LSM sampling method resulted in a higher frequency of detects (number of detections) than other methods. The average frequency of detected congeners for the LSM sampling method over all events was: total – 38% (1.5 detects out of 4 compounds) and dissolved – 38% (1.5 detects out of 4 compounds) (Table G-1). No compounds were positively detected in the HSM particulate or LSM particulate sampling methods.
	Where detected in both the HSM and LSM methods, the HSM total concentrations were higher (19%), on average, than the LSM total concentrations (Table G-1).

	Where detected in both the HSM and whole water methods, the HSM total concentrations were lower (-32%), on average, than the whole water concentrations (Table G-1).
	Where detected in both the LSM and whole water methods, the LSM total concentrations were lower (-36%), on average, than the whole water concentrations (Table G-1).
	Where detected in both the HSM and LSM dissolved sampling methods, the HSM dissolved concentrations were higher (19%), on average, than the LSM dissolved concentrations (Table G-1).
Res	sults of the chi-square test indicated the following:
	There was no significant difference in frequency of detection among methods (HSM, LSM, and whole water) (Table G-2).
Tab	oles G-3 through G-7 provide the analytical results and conversions for each method and sampling event
2.8	Cyanide
Sar Eve for	o of the three sample collection and processing methods (HSM and whole water) were evaluated for CN. mples (primary sample and field duplicate sample) were collected for CN analysis during two events: ent #1/Attempt #2 and Event #2/Attempt #2. A summary of the findings of the additional data evaluations CN data are provided below. Data evaluation summaries and analytical results are presented in pendix H.
	The frequency of detection was the same for HSM total and whole water concentrations (100%) (Table H-1).
******	Where detected in both HSM and whole water sampling methods, the HSM total concentrations were lower (-43%), on average, than the whole water concentrations. However, it should be noted that total concentrations in Event #1/Attempt #2 were similar between HSM total and whole water, but whole water concentrations were of a magnitude (approximately 10 times) greater than HSM total in Event #2/Attempt #2 (Table H-1).
CN	was detected in all samples that were analyzed using HSM and whole water sampling methods

Therefore, the chi-square test was not conducted for this compound. As discussed above, CN was not

Tables H-2 through H-4 provide the analytical results and conversions for each method and sampling event.

analyzed for LSM particulate/dissolved samples.

2.9 Volatile Organic Compounds

Two of the three sample collection and processing methods (HSM and whole water) were evaluated for VOCs. VOCs were not analyzed using the LSM method due to the required filtration method that compromises sample integrity for VOCs. Samples (primary sample and field duplicate sample) were collected for VOC analysis during two events: Event #1/Attempt #2 and Event #2/Attempt #1. A summary of the findings of the additional data evaluations for VOC data are provided below. Data evaluation summaries and analytical results are presented in Appendix I.

and	d analytical results are presented in Appendix I.
-	Chlorobenzene was positively identified (above the PQL) during HSM particulate analysis only.
	1,4-Dichlorobenzene was positively identified during HSM particulate, HSM dissolved, and whole water analyses; however, it was only positively detected (above the PQL) during HSM particulate analysis (Table I-1).
Amenda	Concentration comparisons were not performed because VOCs were detected only for HSM particulate analysis.
Re	sults of the chi-square test indicated the following:
	Frequency of detection was not significantly different between methods (HSM and whole water) according to the Fisher Exact Test (Table I-2). As discussed above, VOCs were not analyzed for LSM particulate/dissolved samples.
Tal	oles I-3 through I-5 provide the analytical results and conversions for each method and sampling event.
2.1	0 Total Extractable Petroleum Hydrocarbons
TE eve	o of the three sample collection and processing methods (HSM and whole water) were evaluated for PH. Samples (primary sample and field duplicate sample) were collected for TEPH analysis during two ents: Event #1/Attempt #2 and Event #2/Attempt #2. A summary of the findings of the additional data aluations for TEPH data are provided below. Data evaluation summaries and analytical results are esented in Appendix J.
and an analysis	The frequency of detection was the same (equal) for HSM total and whole water concentrations (100%) (Table J-1).
decision	Where detected in both the HSM and whole water methods, the HSM total concentrations were lower (-55%), on average, than the whole water concentrations (Table J-1).

TEPH was detected in all samples that were analyzed using the HSM and whole water sampling methods. Therefore, the chi-square test was not conducted for this compound. As discussed above, TEPH was not analyzed for LSM particulate/dissolved samples.

Tables J-2 through J-4 provide the analytical results and conversions for each method and sampling event.

3. Summary

This Phase I Report Addendum presents the additional data evaluations conducted for each analytical group and includes detailed statistical analyses to compare the frequency of detections and differences in concentrations between each sampling method used in Phase I (HSM, LSM, and whole water). These additional data evaluations were beyond the scope of the data evaluation criteria defined in the CSO/SWO Investigation QAPP (Tierra 2013) and the results generated by these additional data evaluations provide a more in-depth analysis of the Phase I data than provided in the Phase I Report.

The summary results presented in Section 2 show the observed differences with respect to number of detections and concentrations for each analytical group. HSM is the most sensitive sampling method with respect to the number of detections and provides the best approach for detecting target compounds present in CSO and SWO overflow. Drawing meaningful conclusions regarding the relative concentrations of target compounds observed between sampling methods is more challenging given the observed variability between sampling events/attempts and analytical groups.

4. References

- Agresti, A. 1990. Categorical Data Analysis. John Wiley and Sons. New York.
- Tierra. 2013. Combined Sewer Overflow/Stormwater Outfall Investigation Quality Assurance Project Plan. Lower Passaic River Study Area. Revision 3. September 2013.
- Tierra. 2014. Combined Sewer Overflow/Stormwater Outfall Phase I Evaluation/Recommendation Report, Revision 0. October 2014.
- Tierra. 2016. Combined Sewer Overflow/Stormwater Outfall Phase I Evaluation/Recommendation Report, Revision 1. March 2016.

Appendix A Data Evaluation Summaries and Analytical Results – PCDDs/PCDFs

Table A-1
Summary of Detected Dioxin Congeners by Method and Event
Phase I Report Addendum – Additional Data Evaluation

Analyte (Dioxin)	Event / Attempt		Concenti (pg/L)		Parent a Partio	ulate	licate Sample) Dissolved (pg/L) HSM LSM		Percent Difference for HSM Compared to Other Methods for Total Concentrations (pg/L) When Detected by Both Methods LSM WW		Percent Difference for LSM Compared to WW for Total Concentrations (pg/L) When Detected by Both Methods	Percent Difference for HSM Compared to LSM for Particulate Concentrations (pg/L) When Detected by Both Methods	Percent Difference for HSM Compared to LSM for Dissolved Concentrations (pg/L) When Detected by Both Methods
2,3,7,8-TETRACHLORODIBENZO-P-DIOXIN	All	2	0	0	2	0	0	0	LSIVI	***************************************	Both Methods	Dotti Wethous	Dotti Wethous
1,2,3,7,8-PENTACHLORODIBENZO-P-DIOXIN	All	2	0	0	2	0	0	0					
1,2,3,4,7,8+EXACHLORODIBENZO-P-DIOXIN	All	6	0	0	6	0	0	0					
1,2,3,6,7,8HEXACHLORODIBENZO-P-DIOXIN	All	6	0	2	6	0	1	0		-22%			
1,2,3,7,8,9 HEXACHLORODIBENZO-P-DIOXIN	All	6	0	0	6	0	0	0					
1,2,3,4,6,7,8HEPTACHLORODIBENZO-P-DIOXIN	All	6	6	6	6	6	6	6	18%	4%	-11%	-48%	491%
OCTACHLORODIBENZO-P-DIOXIN	All	6	6	6	6	6	6	6	7%	-17%	-22%	-43%	564%
2,3,7,8-TETRACHLORODIBENZOFURAN	All	6	0	0	6	0	0	0			·		·
1,2,3,7,8-PENTACHLORODIBENZOFURAN	All	2	0	0	2	0	0	0					
2,3,4,7,8-PENTACHLORODIBENZOFURAN	All	3	0	0	3	0	0	0					
1,2,3,4,7,8HEXACHLORODIBENZOFURAN	All	4	0	0	4	0	0	0					
1,2,3,6,7,8HEXACHLORODIBENZOFURAN	All	6	0	0	6	0	0	0					
2,3,4,6,7,&HEXACHLORODIBENZOFURAN	All	6	0	0	6	0	0	0					
1,2,3,7,8,9HEXACHLORODIBENZOFURAN	All	0	0	0	0	0	0	0					
1,2,3,4,6,7,8HEPTACHLORODIBENZOFURAN	All	6	4	4	6	4	6	3	28%	5%	-17%	-51%	403%
1,2,3,4,7,8,9HEPTACHLORODIBENZOFURAN	All	6	0	1	6	0	0	0		-68%			
OCTACHLORODIBENZOFURAN	All	6	4	4	6	4	6	2	48%	8%	-25%	-50%	543%
Summary													
17 Congeners	1/1	13.5	4	4	13.5	4	4.5	4	9%	13%	4%	-92%	798%
17 Congeners	1/3	15	4	4.5	15	4	4	2.5	47%	6%	-28%	-4%	260%
17 Congeners	2/2	11	2	3	11	2	4	2	0%	-51%	-43%	-47%	268%
17 Congeners	1/3 and 2/2 Only	13	3	3.8	13	3	4	2.3	32%	-20%	-33%	-18%	263%
17 Congeners	All	13.2	3	3.8	13.2	3	4.2	2.8	23%	-10%	-18%	-48%	501%
Percent of 17 Detected Congeners	All	77%	20%	23%	77%	20%	25%	17%					

Summary:

HSM has a higher frequency of detection (number of detections) for total (77%), particulate (77%), and dissolved (25%) concentrations.

Where detected in both methods, HSM total concentrations are on average 23% greater than LSM total concentrations.

Where detected in both methods, HSM total concentrations are slightly lower on average (-10%) than WW concentrations; however, there is great variability among events.

Where detected in both methods, LSM total concentrations are on average lower than WW concentrations (-18%).

Where detected in both methods, HSM particulate concentrations (pg/L) are on average lower than LSM particulate concentrations (pg/L) (-48%).

Where detected in both methods, HSM dissolved concentrations are on average much higher than LSM dissolved concentrations (501%).

<u>Abbreviations</u>

pg/L = picograms per liter % = percent HSM = high solids mass LSM = low solids mass WW = whole water

Table A-2
Statistical Comparison of the Number of Detected Dioxin Congeners by Method and Event
Phase I Report Addendum – Additional Data Evaluation

	Number	of Detectio	ns (Total								
	Wate	r Concentra	ation)	Maximum Possible	Chi-Square Test (p-value) ²						
Event	HSM LSM WW		Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW					
1/1	27 8 8		34	<0.001	<0.001	1					
1/3	30	8	9	34	<0.001	<0.001	0.78				
2/2	22	4	6	34	<0.001	<0.001	0.49				
All	79	20	23	102	<0.001	<0.001	0.61				

Notes

Conclusions

The HSM method is better than both other methods with respect to the number of detected congeners.

The LSM and WW methods are similar with respect to the number of detected congeners.

Abbreviations

HSM = high solids mass

LSM = low solids mass

WW = whole water

¹ Total number of detections for event includes 2 duplicates and 17 congeners.

² A p-value less than 0.05 is considered significant and is shaded indicating that the number of detects is significantly different between methods.

Table A-3 HSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

		HSM Particulate Sample Collection																
		Eve	ent 2 Attempt 2	2 (12-7-13 P	R135)			Eve	ent 1 Attempt	3 (4-30-14 PF	R146)			Ev	ent 1 Attempt	1 (6-10-13 P	R106)	
	PR	1CSOCLYHF	P-02B	PR1HPDUP-02B			PF	R1CSOCLYHI	P-01C	Р	R1HPDUP	01C	PR	1CSOCLYH	P-01A	PR1HPDUP-01A		
Wet weight (gram)		13.8			15.4			19.9			19.2			17.3		16.9		
% Solids		36.3			36.4			50.2			52			29.5			30.1	
Compound Identified	Weight gram (dry)	Sample Result pg/g	Converted Sample Result pg/L	Weight gram (dry)	Sample Result pg/g	Converted Sample Result pg/L	Weight gram (dry)	Sample Result pg/g	Converted Sample Result pg/L									
2,3,7,8-TETRACHLORODIBENZOP-DIOXIN	5.01		0	5.61		0	9.99		0	9.98		0	5.10	2.36	0.0120	5.09	9.15	0.0471
1,2,3,7,8-PENTACHLORODIBENZO-P-DIOXIN	5.01		0	5.61		0	9.99	4.56	0.252	9.98	4.69	0.268	5.10		0	5.09		0
1,2,3,4,7,8-HEXACHLORODIBENZO-P-DIOXIN	5.01	6.32	0.204	5.61	6.16	0.199	9.99	9.01	0.498	9.98	9.24	0.529	5.10	5.96	0.0302	5.09	5.72	0.0295
1,2,3,6,7,8-HEXACHLORODIBENZO-P-DIOXIN	5.01	21.1	0.680	5.61	19.8	0.640	9.99	24.4	1.35	9.98	25	1.43	5.10	21.4	0.109	5.09	21.2	0.109
1,2,3,7,8,9-HEXACHLORODIBENZO-P-DIOXIN	5.01	15.2	0.490	5.61	14.2	0.459	9.99	17.5	0.968	9.98	21	1.20	5.10	15.3	0.0776	5.09	15.3	0.0788
1,2,3,4,6,7,8-HEPTACHLORODIBENZO-P-DIOXIN	5.01	700	22.6	5.61	636	20.6	9.99	746	41.2	9.98	818	46.8	5.10	672	3.41	5.09	621	3.20
OCTACHLORODIBENZO-P-DIOXIN	5.01	9590	309	5.61	9560	309	9.99	12000	663	9.98	11600	664	5.10	9480	48.1	5.09	8960	46.2
2,3,7,8-TETRACHLORODIBENZOFURAN	5.01	3.82	0.123	5.61	2.88	0.0931	9.99	3.85	0.213	9.98	3.6	0.206	5.10	4.76	0.0241	5.09	4.9	0.0252
1,2,3,7,8-PENTACHLORODIBENZOFURAN	5.01		0	5.61		0	9.99	3.53	0.195	9.98	3.22	0.184	5.10		0	5.09		0
2,3,4,7,8-PENTACHLORODIBENZOFURAN	5.01		0	5.61		0	9.99	4.77	0.264	9.98	4.21	0.241	5.10	0	0	5.09	5.26	0.0271
1,2,3,4,7,8-HEXACHLORODIBENZOFURAN	5.01		0	5.61		0	9.99	14.9	0.824	9.98	14.4	0.824	5.10	20.9	0.106	5.09	31.5	0.162
1,2,3,6,7,8-HEXACHLORODIBENZOFURAN	5.01	11.7	0.377	5.61	11.1	0.359	9.99	13.9	0.769	9.98	14.2	0.813	5.10	15.4	0.0781	5.09	18.2	0.0938
2,3,4,6,7,8-HEXACHLORODIBENZOFURAN	5.01	10.5	0.338	5.61	7.89	0.255	9.99	9.96	0.551	9.98	10.5	0.601	5.10	19	0.0964	5.09	20.9	0.108
1,2,3,7,8,9 HEXACHLORODIBENZOFURAN	5.01		0	5.61		0	9.99		0	9.98		0	5.10		0	5.09		0
1,2,3,4,6,7,8-HEPTACHLORODIBENZOFURAN	5.01	205	6.605	5.61	197	6.37	9.99	253	14.0	9.98	247	14.1	5.10	245	1.24	5.09	271	1.40
1,2,3,4,7,8,9-HEPTACHLORODIBENZOFURAN	5.01	13.3	0.429	5.61	12.5	0.404	9.99	13.8	0.763	9.98	14.4	0.824	5.10	16.4	0.0832	5.09	18.7	0.0963
OCTACHLORODIBENZOFURAN	5.01	444	14.3	5.61	458	14.8	9.99	488	27.0	9.98	469	26.8	5.10	486	2.46	5.09	549	2.83

<u>Abbreviations</u>

pg/L = picograms per liter pg/g = picograms per gram

Table A-4
HSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

	HSM Dissolved Sample Collection														
	Even	t 2 Attempt	2 (12-7-13 P	R138)	Even	t 1 Attempt	3 (4-30-14 P	R147)	Even	t 1 Attempt	1 (6-10-13 PI	R107)			
	PR1CSOC	LYHD-02B	PR1HDI	OUP-02B	PR1CSOC	LYHD-01C	PR1HDI	OUP-01C	PR1CSOCLYHD-01A		PR1HDE	OUP-01A			
Compound Identified	Volume Liters	Sample Result pg/L	Volume Liters	Sample Result pg/L	Volume Liters	Sample Result pg/L	Volume Liters	Sample Result pg/L	Volume Liters	Sample Result pg/L	Volume Liters	Sample Result pg/L			
2,3,7,8-TETRACHLORODIBENZO-P-DIOXIN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,7,8-PENTACHLORODIBENZO-P-DIOXIN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,4,7,8-HEXACHLORODIBENZO-P-DIOXIN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,6,7,8-HEXACHLORODIBENZO-P-DIOXIN	9.66		9.84		9.74		10.0		9.91	0	9.88	4.63			
1,2,3,7,8,9-HEXACHLORODIBENZO-P-DIOXIN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,4,6,7,8-HEPTACHLORODIBENZO-P-DIOXIN	9.66	38.5	9.84	30.5	9.74	31.3	10.0	29.3	9.91	32.6	9.88	116			
OCTACHLORODIBENZO-P-DIOXIN	9.66	338	9.84	199	9.74	226	10.0	269	9.91	365	9.88	720			
2,3,7,8-TETRACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,7,8-PENTACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
2,3,4,7,8-PENTACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,4,7,8-HEXACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,6,7,8-HEXACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
2,3,4,6,7,8-HEXACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,7,8,9-HEXACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
1,2,3,4,6,7,8-HEPTACHLORODIBENZOFURAN	9.66	17.3	9.84	13.4	9.74	15.3	10.0	13.0	9.91	16.6	9.88	17.6			
1,2,3,4,7,8,9-HEPTACHLORODIBENZOFURAN	9.66		9.84		9.74		10.0		9.91		9.88				
OCTACHLORODIBENZOFURAN	9.66	42.3	9.84	32.5	9.74	26.8	10.0	23.1	9.91	37.0	9.88	39.8			

Abbreviations

pg/L = picograms per liter

Table A-5
LSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

	LSM Particulate Sample Collection																			
		Even	t 2 Attempt 2	2 (12-7-13 P	R140)			Even	t 1 Attempt 3	3 (4-30-14 PI	R149)		Event 1 Attempt 1 (6-10-13 PR109)							
	PR:	LCSOCLYLP-	02B	P	R1LPDUP-02	2B	PR:	LCSOCLYLP	01C	P	R1LPDUP-01	LC	PR1CSOCLYLP-01A			PR1LPDUP-01A				
Total Liters Filtered (L)		9.476			9.491			9.663			10.103		10.035			9.713				
			Converted			Converted			Converted			Converted			Converted			Converted		
		Sample	Sample		Sample	Sample		Sample	Sample		Sample	Sample		Sample	Sample		Sample	Sample		
Compound Identified	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result		
	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L		
2,3,7,8-TETRACHLORODIBENZO-P-DIOXIN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,7,8-PENTACHLORODIBENZO-P-DIOXIN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,4,7,8-HEXACHLORODIBENZO-P-DIOXIN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,6,7,8-HEXACHLORODIBENZO-P-DIOXIN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,7,8,9-HEXACHLORODIBENZO-P-DIOXIN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,4,6,7,8-HEPTACHLORODIBENZO-P-DIOXIN	0.0796	4920	41.3	0.128	3160	42.6	0.0773	3750	30.0	0.0808	7400	59.2	0.371	1940	71.7	0.612	845	53.2		
OCTACHLORODIBENZO-P-DIOXIN	0.0796	64000	538	0.128	43100	581	0.0773	45500	364	0.0808	109000	872	0.371	15700	580	0.612	8560	539		
2,3,7,8-TETRACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,7,8-PENTACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
2,3,4,7,8-PENTACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,4,7,8-HEXACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,6,7,8-HEXACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
2,3,4,6,7,8-HEXACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,7,8,9-HEXACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
1,2,3,4,6,7,8-HEPTACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773	1760	14.1	0.0808	2230	17.8	0.371	396	14.6	0.612	215	13.5		
1,2,3,4,7,8,9-HEPTACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773		0	0.0808		0	0.371		0	0.612		0		
OCTACHLORODIBENZOFURAN	0.0796		0	0.128		0	0.0773	3280	26.2	0.0808	4070	32.6	0.371	790	29.2	0.612	432	27.2		

<u>Abbreviations</u>

pg/L = picograms per liter pg/g = picograms per gram

Table A-6
LSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

	LSM Dissolved Sample Collection											
	Even	t 2 Attempt 2	2 (12-7-13 PR	141)	Even	t 1 Attempt 3	(4-30-14 PR	150)	Event 1 Attempt 1 (6-10-13 PR110)			
	PR1CSOCLYLD-02B		PR1LDDUP-02B		PR1CSOC	CLYLD-01C	PR1LDDUP-01C		PR1CSOCLYLD-01A		PR1LDD	UP-01A
	Sample		Sample			Sample		Sample		Sample		Sample
	Volume	Result	Volume	Result	Volume	Result	Volume	Result	Volume	Result	Volume	Result
Compound Identified	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L
2,3,7,8-TETRACHLORODIBENZO-P-DIOXIN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,7,8-PENTACHLORODIBENZO-P-DIOXIN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,4,7,8-HEXACHLORODIBENZO-P-DIOXIN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,6,7,8-HEXACHLORODIBENZO-P-DIOXIN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,7,8,9-HEXACHLORODIBENZO-P-DIOXIN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,4,6,7,8-HEPTACHLORODIBENZO-P-DIOXIN	9.93	11.0	9.72	8.92	9.90	13.0	9.99	10.4	9.79	6.33	9.71	6.41
OCTACHLORODIBENZO-P-DIOXIN	9.93	73.2	9.72	64.7	9.90	74.90	9.99	72.8	9.79	41.7	9.71	44.0
2,3,7,8-TETRACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,7,8-PENTACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
2,3,4,7,8-PENTACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,4,7,8-HEXACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,6,7,8-HEXACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
2,3,4,6,7,8-HEXACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,7,8,9-HEXACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
1,2,3,4,6,7,8-HEPTACHLORODIBENZOFURAN	9.93		9.72		9.90	5.81	9.99	0	9.79	3.40	9.71	3.20
1,2,3,4,7,8,9-HEPTACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79		9.71	
OCTACHLORODIBENZOFURAN	9.93		9.72		9.90		9.99		9.79	6.05	9.71	5.9

Abbreviations

pg/L = picograms per liter

Table A-7
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

	Whole Water Sample Collection												
	Even	t 2 Attempt 2	2 (12-7-13 PI	R134)	Event	t 1 Attempt 3	3 (4-30-14 PI	R145)	Event 1 Attempt 1 (6-10-13 PR105)				
	PR1CSOCI	.YWW-02B	PR1WW	DUP-02B	PR1CSOCLYWW-01C		PR1WWDUP-01C		PR1CSOCLYWW-01A		PR1WWDUP-01A		
		Sample		Sample		Sample		Sample		Sample		Sample	
Compound Identified	Volume	Result	Volume	Result	Volume	Result	Volume	Result	Volume	Result	Volume	Result	
	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L	
2,3,7,8-TETRACHLORODIBENZO-P-DIOXIN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,7,8-PENTACHLORODIBENZO-P-DIOXIN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,4,7,8-HEXACHLORODIBENZO-P-DIOXIN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,6,7,8-HEXACHLORODIBENZO-P-DIOXIN	9.73		9.63	2.76	9.78	2.58	9.67		7.23		9.5		
1,2,3,7,8,9-HEXACHLORODIBENZO-P-DIOXIN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,4,6,7,8-HEPTACHLORODIBENZO-P-DIOXIN	9.73	84.3	9.63	87.4	9.78	81.5	9.67	71.1	7.23	62.1	9.5	41.3	
OCTACHLORODIBENZO-P-DIOXIN	9.73	1090	9.63	1230	9.78	1060	9.67	821	7.23	715	9.5	429	
2,3,7,8-TETRACHLORODIBENZOFURAN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,7,8-PENTACHLORODIBENZOFURAN	9.73		9.63		9.78		9.67		7.23		9.5		
2,3,4,7,8-PENTACHLORODIBENZOFURAN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,4,7,8-HEXACHLORODIBENZOFURAN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,6,7,8-HEXACHLORODIBENZOFURAN	9.73		9.63		9.78		9.67		7.23		9.5		
2,3,4,6,7,8-HEXACHLORODIBENZOFURAN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,7,8,9-HEXACHLORODIBENZOFURAN	9.73		9.63		9.78		9.67		7.23		9.5		
1,2,3,4,6,7,8-HEPTACHLORODIBENZOFURAN	9.73		9.63		9.78	29.1	9.67	20.2	7.23	18	9.5	20.5	
1,2,3,4,7,8,9-HEPTACHLORODIBENZOFURAN	9.73		9.63	2.61	9.78		9.67		7.23		9.5		
OCTACHLORODIBENZOFURAN	9.73		9.63		9.78	53.7	9.67	38.0	7.23	36.6	9.5	43.2	

Abbreviations

pg/L = picograms per liter

Appendix B Data Evaluation Summaries and Analytical Results – PCB Congeners

		<u> </u>								erence for HSM Other Methods for	Percent Difference	Percent Difference for	Percent Difference for
Analyte (PCBs)			er of Dete						Total Conce	ntrations (pg/L)	for LSM Compared to WW for Total	HSM Compared to LSM for Particulate	HSM Compared to LSM
		Total (Concentr (pg/L)	ncentrations og/L)		ulates z/L)	tes Dissolv (pg/			ected by Both thods	Concentrations (pg/L)	Concentrations (pg/L)	for Dissolved Concentrations (pg/L)
	Event/ Attempt	HSM	LSM	ww	HSM	LSM	HSM	LSM	LSM	ww	When Detected by Both Methods	When Detected by Both Methods	When Detected by Both Methods
PCB-1 PCB-2	All	0	0	4 0	2	0	0	0	66%	47%	-11%		25%
PCB-3	All	0	0	0	0	0	0	0	F70/	710/	F.00/	010/	
PCB-4/10 PCB-5/8	All	6 4	3	2	6 4	0	0	0	-57%	-71% -95%	-50%	-81%	
PCB-6 PCB-7/9	All All	4 0	0	3	4 0	0	2	0		57%			
PCB-11	All	2	0	2	2	0	0	0					
PCB-12/13 PCB-14	All	0	0	0	0	0	0	0					
PCB-15	All	5	0	2	5	0	0	0		-93%			
PCB-16/32 PCB-17	All	6 6	2	3	6	2	0	0	-89% -88%	-25% -50%	-45% -38%	-89% -88%	
PCB-18	All	6	2	2	6	2	0	0	-89%	-94%	-43%	-89%	
PCB-19 PCB-20/21/33	All	6 5	2	5 2	6 5	2	0	0	-89% -93%	-39% -96%	-51% -45%	-89% -93%	
PCB-22 PCB-23	All	6 0	2	2	6 0	2	0	0	-94%	-96%	-38%	-94%	
PCB-24/27	All	4	0	2	4	0	0	0		-22%			
PCB-25 PCB-26	All All	6 6	1 2	5 3	6 6	2	2	0	172% -93%	152% -17%	-61% -37%	-89% -93%	
PCB-28	All	6	2	3	6	2	0	0	-93% -92%	-12%	-38%	-92%	
PCB-29 PCB-30	All All	0	0	0	0	0	0	0					
PCB-31	All	6	2	2	6	2	0	0	-93%	-96%	-33%	-93%	
PCB-34 PCB-35	All All	0 5	0	0	0 5	0	0	0	118%	-50%	-56%	118%	
PCB-36	All	0	0	0	0	0	0	0	118%		-50%	118%	
PCB-37 PCB-38	All All	5 0	2	2	5 0	2	0	0	-92%	-95%	-39%	-92%	
PCB-39	All	0	0	0	0	0	0	0					
PCB-40 PCB-41/64/71/72	All All	6 6	3	3 4	6 6	3	0	0	-3% 51%	11% 1%	-10% -33%	-3% 7%	
PCB-42/59	All	6	3	4	6	3	0	0	13%	-36%	-36%	13%	
PCB-43/49 PCB-44	All All	6 6	3	4	6 6	3	2	0	48% 28%	-5% -19%	-34% -32%	3% 8%	
PCB-45	All	6	3	3	6	3	0	0	-17%	-12%	-17%	-17%	
PCB-46 PCB-47	All All	6 3	2	4 3	6 3	2	0	0	-1% 136%	30% 98%	-41% -39%	⁻ 1% 136%	
PCB-48/75	All	6	3	5	6	3	0	0	-29%	-10%	-9%	-29%	
PCB-50 PCB-51	All	0 5	1 2	2	0 5	1 2	0	0	26%	-9%	-65% -29%	26%	
PCB-52/69	All	6	3	4	6	3	2	0	36%	-13%	-34%	-4%	
PCB-53 PCB-54	All	6 1	3	4	6 1	3	0	0	39%	-28%	-43%	9%	
PCB-55	All	1	0	0	1	0	0	0					
PCB-56/60 PCB-57	All	6 1	3	3	6 1	3	0	0	29%	35%	-5%	-4%	
PCB-58	All	0	0	0	0	0	0	0					
PCB-61/70 PCB-62	All	6	3	5	6	3	2 0	0	28%	30%	-31%	-6%	
PCB-63	All	4	1	2	4	1	0	0	17%	-37%	2%	17%	
PCB-65 PCB-67	All	2	0	0	2	0	0	0					
PCB-68	All	0	0	0	0	0	0	0					
PCB-73 PCB-74	All	0 6	3	0 4	0 6	3	2	0	16%	-4%	-13%	-11%	120%
PCB-76/66	All	6	3	5	6	3	2	0	39%	53%	-30%	4%	
PCB-77 PCB-78	All	4 0	3	3	0	3	0	0	19%	16%	-13%	19%	
PCB-79	All	2	0	0	2	0	0	0					
PCB-80 PCB-81	All	0	0	0	0	0	0	0					
PCB-82	All	6	4	6	6	4	4	2	16%	-3%	-18%	-10%	66%
PCB-83 PCB-84/92	All	6	0 4	0 6	0 6	0 4	2	0	15%	-11%	-30%	-13%	
PCB-85/116	All	6	6	6	6	4	6	4	199%	21%	-34%	-7%	96%
PCB-86 PCB-87/117/125	All	6	5	0 6	6	5	3	0	70%	-8%	-43%	43%	
PCB-88/91 PCB-89	All	6 1	4	6	6	4	4	2	11%	1%	-16%	-17%	63%
PCB-90/101	All	6	4	6	6	4	5	0	26%	13%	-31%	-13%	
PCB-93 PCB-94	All All	0	0	0	0	0	0	0					
PCB-95/98/102	All	6	4	6	6	4	0	0	-18%	-28%	-32%	-18%	
PCB-96 PCB-97	All All	0 6	0 4	0 6	0 6	0 4	0	0	22%	-12%	-33%	-7%	
PCB-99	All	6	4	6	6	4	6	0	44%	24%	-30%	-6%	
PCB-100 PCB-103	All All	0	0	0	0	0	0	0					
PCB-104	All	0	0	0	0	0	0	0					
PCB-105 PCB-106/118	All All	6 6	5 5	6 6	6 6	5 5	4 5	0	77% 98%	-1% 9%	-43% -40%	44% 46%	
PCB-107/109	All	6	4	5	6	4	3	0	5%	82%	-14%	-6%	
PCB-108/112 PCB-110	All All	6 6	3 5	6 6	6 6	3 5	2 4	0	47% 102%	-14% 0%	-42% -42%	15% 71%	
PCB-111/115	All	4	2	1	4	2	0	0	-87%	-87%	4%	-87%	
PCB-113 PCB-114	All All	0	0 2	0 2	0	0 2	0	0	-79%	-85%	-16%	-79%	
PCB-119	All	4	0	1	4	0	0	0	73/0	-87%	1076	1376	
PCB-120 PCB-121	All All	0	0	0	0	0	0	0					
PCB-122	All	1	0	1	1	0	0	0					
PCB-123 PCB-124	All All	4 6	1 3	0 6	4 6	1 3	0	0	-86% 31%	-27%	-40%	-86% 31%	
PCB-126	All	3	0	0	3	0	0	0	31/6	21/0	40%	51%	
PCB-127	All	0	0	0	0	0	0	0					

		Numbe	er of Dete	ections (Parent	and Dup	olicate S	ample)	Percent Difference for HSM Compared to Other Methods for Total Concentrations (pg/L)		Percent Difference for LSM Compared to	Percent Difference for HSM Compared to LSM	Percent Difference for HSM Compared to LSM
Analyte (PCBs)		Total (Concentr (pg/L)	rations		culates g/L)		olved g/L)	When Dete	cted by Both thods	WW for Total Concentrations (pg/L) When Detected by	for Particulate	for Dissolved Concentrations (pg/L) When Detected by Both
	Event/ Attempt	HSM	LSM	ww	нѕм	LSM	HSM	LSM	LSM	ww	Both Methods	Methods	Methods
PCB-128/162	All	6	5	6	6	5	6	2	92%	15%	-36%	46%	63%
PCB-129 PCB-130	All	6 6	4	6 6	6 6	4	3	0	19% 9%	-8% 1%	-39% -32%	-7% -1%	
PCB-131	All	0	0	0	0	0	0	0	370	170	32/0	170	
PCB-132/161	All	6	5	6	6	5	4	0	102%	7%	-42%	50%	
PCB-133/142	All	6	3	5	6	3	0	0	19%	6%	-41%	19%	
PCB-134/143 PCB-135	All	6 6	4	6 6	6 6	4	2 6	2	31% 5%	-12% 15%	-40% -14%	1% -23%	53%
PCB-136	All	6	4	5	6	4	4	1	11%	36%	-23%	-24%	221%
PCB-137	All	6	4	6	6	4	2	0	-20%	5%	-14%	-20%	
PCB-138/163/164	All	6	5	6	6	5	6	0	113%	11%	-44%	47%	540
PCB-139/149 PCB-140	All	6 0	4 0	6 0	6 0	0	4 0	2	4%	1%	-15%	-22%	61%
PCB-141	All	6	4	6	6	4	2	0	15%	-16%	-36%	-9%	
PCB-144	All	6	4	5	6	4	0	0	-30%	18%	-24%	-30%	
PCB-145	All	0	0	0	0	0	0	0					
PCB-146/165	All	6	4	6	6	4	2	0	18%	-13%	-35%	-7%	
PCB-147 PCB-148	All	5 0	0	0	5	0	0	0		-87%			
PCB-150	All	0	0	0	0	0	0	0					
PCB-151	All	6	4	4	6	4	1	2	-33%	-34%	-17%	-29%	-16%
PCB-152	All	0	0	0	0	0	0	0	0.004	A	2011	6	
PCB-153 PCB-154	All	6 1	4 0	6	6 1	0	4 0	0	36%	-6%	-36%	-9%	
PCB-155	All	0	0	0	0	0	0	0					
PCB-156	All	6	4	6	6	4	4	0	39%	-6%	-37%	0%	
PCB-157	All	6	2	4	6	2	0	0	49%	34%	-38%	49%	
PCB-158/160	All	6	4	6	6	4	2	0	20%	-11%	-37%	-4%	
PCB-159 PCB-166	All	0 2	0	0	2	0	0	0					
PCB-167	All	6	3	6	6	3	0	0	48%	-23%	-49%	48%	
PCB-168	All	0	0	0	0	0	0	0					
PCB-169	All	0	0	0	0	0	0	0					
PCB-170	All	6	5 4	6	6	5	4	2	53%	-7% -27%	-37% -30%	34%	67%
PCB-171 PCB-172	All	6 6	3	6 6	6 6	3	0	0	1% -4%	-27% -29%	-39% -50%	-12% -4%	
PCB-173	All	1	0	0	1	0	0	0	470	2370	30/0	470	
PCB-174	All	6	4	4	6	4	4	2	13%	-24%	-34%	-13%	58%
PCB-175	All	2	0	0	2	0	0	0					
PCB-176 PCB-177	All	6 6	2	3 6	6 6	2	0 4	0 2	2% 14%	-46% -11%	-44% -34%	2% -11%	73%
PCB-178	All	6	3	5	6	3	0	0	-18%	-14%	-28%	-18%	7 3 7 0
PCB-179	All	6	3	4	6	3	0	0	-1%	-46%	-45%	-1%	
PCB-180	All	6	4	4	6	4	0	0	-13%	-46%	-37%	-13%	
PCB-181	All	0	0	0	0	0	0	0	2504	2.424		2.21	
PCB-182/187 PCB-183	All	6 6	4	4	6 6	4	2	2	-25% -18%	-34% -32%	-24% -26%	-26% -19%	57% 56%
PCB-184	All	0	0	0	0	0	0	0	1870	32/0	20/0	1370	30/0
PCB-185	All	6	2	3	6	2	0	0	6%	-52%	-49%	6%	
PCB-186	All	0	0	0	0	0	0	0					
PCB-188 PCB-189	All	0 3	0	0	3	0	0	0					
PCB-189	All	6	4	4	6	4	0	0	-14%	-51%	-46%	-14%	
PCB-191	All	1	0	0	1	0	0	0	1170		1370	1470	
PCB-192	All	0	0	0	0	0	0	0					
PCB-193	All	6	2	3	6	2	0	0	25%	-26%	-55% -27%	25%	
PCB-194 PCB-195	All	6 5	4	4 6	6 5	3	0	0	-4% 8%	-40% -40%	-37% -47%	⁻ 17%	
PCB-196/203	All	6	4	4	6	4	2	2	-35%	-35%	-10%	-38%	58%
PCB-197	All	0	0	0	0	0	0	0		_			
PCB-198	All	0	0	0	0	0	0	0					
PCB-199 PCB-200	All	6 5	4	6 2	6 5	4	0	0	-34% 26%	-9% -30%	⁻ 20% 7%	-36% 26%	57%
PCB-200 PCB-201	All	6	1	2	6	1	0	0	46%	-30%	-16%	26% 46%	
PCB-202	All	6	2	4	6	2	0	0	-14%	15%	-36%	-14%	
PCB-204	All	0	0	0	0	0	0	0					
PCB-205	All	0	0	0	0	0	0	0					
PCB-206 PCB-207	All	6 1	2	4	6 1	0	0	0	25%	-42%	-51%	25%	
PCB-207 PCB-208	All	6	2	3	6	2	0	0	32%	-32%	-37%	32%	
PCB-209	All	4	2	1	4	2	0	0	-82%	-85%	-15%	-82%	
Total PCBs	All	591	298	430	591	292	151	37	412%	28%	-51%	375%	5352%
Summary													
168 Congeners/Coelutions	1/1	98	80.5	92	98	81	39	1.0	60%	-76% 20%	-36% -37%	-87% -70%	120%
168 Congeners/Coelutions 168 Congeners/Coelutions	1/3 2/2	96 102	63 6	79 44.5	96 102	62 4.0	24 13.5	16 2.0	82% 221%	29% 52%	-27% -54%	79% 148%	67% 76%
168 Congeners/Coelutions	1/3 and 2/2 Only	99	34	62	99	33	19	8.8	98%	37%	-30%	85%	68%
168 Congeners/Coelutions	All	99	50	72	99	49	25	6.2	19%	-10%	-33%	-2%	
Percent of 168 Detected Congeners	All	59%	30%	43%	59%	29%	15%	4%					

Conclusions

HSM has a higher frequency of detection for both total (59%), particulate (59%), and dissolved (15%) concentrations.

Where detected in both methods, HSM total concentrations are on average 19% greater than total LSM concentrations; however, there is large variability among events.

Where detected in both methods, HSM total concentrations are slightly lower on average (-10%) than WW concentrations; however, there is great variability among events.

Where detected in both methods, LSM total concentrations are on average lower than WW concentrations (-33%). Where detected in both methods, HSM particulate concentrations are on average slightly lower than LSM particulate concentrations (-2%).

Where detected in both methods, HSM particulate concentrations are on average slightly lower than LSM particulate concentrations. Where detected in both methods, HSM dissolved concentrations are on average 71% greater than LSM dissolved concentrations.

Abbreviations

pg/L = picograms per liter % = percent HSM = high solids mass LSM = low solids mass WW = whole water

Table B-2
Statistical Comparison of the Number of Detected PCB Congeners by Method and Event
Phase I Report Addendum – Additional Data Evaluation

	Number	of Detectio	ns (Total				,
	Wate	r Concentra	ation)	Maximum Possible	Chi-S	quare Test (p-va	lue)²
Event	HSM LSM WW		Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW	
1/1	196	161	184	336	0.0068	0.350	0.070
1/3	191	125	157	336	<0.001	0.009	0.012
2/1	204	12	89	336	<0.001	<0.001	<0.001
All	591	298	430	1008	<0.001	<0.001	<0.001

Notes

Conclusions

The HSM method is better than the LSM method for all events with respect to the number of detected congeners/co-elutions.

The HSM method is better than the WW method for all events with respect to the number of detected congeners/co-elutions; however the difference for event 1/1 is not statistically significant.

The WW method is better than the LSM method for all events with respect to the number of detected congeners/co-elutions.

Abbreviations

HSM = high solids mass

LSM = low solids mass

¹ Total number of detections for event includes 2 duplicates and 168 congeners/co-elutions.

² A p-value less than 0.05 is considered significant and is shaded indicating that the number of detects is significantly different

Table B-3
HSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

								HSM	Particulate :	Sample Colle	ection							
			Event 2 Attemp	ot 2 (12 -7 PR1:	35)				vent 1 Attemp	•				Ev	ent 1 Attempt	1 (6-10-13 PR	106)	-
	P	R1CSOCLYHP		_ `	PR1HPDUP-02	B	P	R1CSOCLYHP-0	•	_ `	PR1HPDUP-01	c	Р	R1CSOCLYHP 0	•	T	PR1HPDUP-01	Δ
Wet weight (gram)		13.9		1	15.6		 	5.85		†	5.8		<u> </u>	20.4		 	19.9	-
% solids		36.4		1	32.9		 	50.2		†	52.0		1	29.5		1	30.1	
,	- 	1	T	1	1		 	T	1	†	1		1	1		1	1	
Compound Identified	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	: Weight gram (dry)	Sample Result	Converted Sample Result pg/L	: Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	: Weight gram (dry)	Sample Result	Converted Sample Result pg/L
PCB-1	5.06	204	7 P8/L	5.13	192	6	2.94	P5/5	0	3.02	P5/5	0	6.02	P5/5	0	5.99	P5/5	0
PCB-2	5.06		0	5.13	192	0	2.94		0	3.02		0	6.02		0	5.99	+	0
PCB-3	5.06	;	0	5.13		0	2.94	+	0	3.02		0	6.02		0	5.99		0
PCB-4/10	5.06	915	30	5.13	1,080	32	2.94	1,550	86	3.02	1,420	81	6.02	870	4.38	5.99	804	4.13
PCB-5/8	5.06		0	5.13	1,000	0	2.94	2,190	121	3.02	1,970	113	6.02	1,340	6.75	5.99	1,270	6.53
PCB-6	5.06	446	14	5.13	639	19	2.94	810	45	3.02	806	46	6.02		0	5.99		0
PCB-7/9	5.06	;	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-11	5.06	;	0	5.13		0	2.94	5,120	283	3.02	4,130	237	6.02		0	5.99	1	0
PCB-12/13	5.06	<u> </u>	0	5.13		0	2.94		0	3.02	İ	0	6.02		0	5.99	1	0
PCB-14	5.06	:	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-15	5.06		0	5.13	1,430	42	2.94	779	43	3.02	819	47	6.02	783	3.94	5.99	706	3.63
PCB-16/32	5.06	1,840	59	5.13	2,250	66	2.94	2,920	162	3.02	3,680	211	6.02	2,260	11.4	5.99	2,180	11.2
PCB-17	5.06	1,250	40	5.13	1,670	49	2.94	2,450	136	3.02	3,360	193	6.02	1,470	7.40	5.99	1,400	7.19
PCB-18	5.06	2,590	84	5.13	2,970	87	2.94	2,820	156	3.02	3,560	204	6.02	2,890	14.6	5.99	2,830	14.5
PCB-19	5.06	420	14	5.13	564	16	2.94	827	46	3.02	933	54	6.02	568	2.86	5.99	581	2.99
PCB-20/21/33	5.06	5	0	5.13	2,230	65	2.94	1,670	92	3.02	1,170	67	6.02	1,130	5.69	5.99	1,050	5.39
PCB-22	5.06	1,140	37	5.13	1,960	57	2.94	1,710	95	3.02	1,100	63	6.02	912	4.59	5.99	679	3.49
PCB-23	5.06	i	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-24/27	5.06		0	5.13		0	2.94	467	26	3.02	605	35	6.02	315	1.59	5.99	305	1.57
PCB-25	5.06	480	16	5.13	4,100	120	2.94	919	51	3.02	1,060	61	6.02	369	1.86	5.99	344	1.77
PCB-26	5.06	701	23	5.13	2,680	78	2.94	1,080	60	3.02	950	54	6.02	608	3.06	5.99	446	2.29
PCB-28	5.06	3,310	107	5.13	15,100	441	2.94	5,920	328	3.02	4,500	258	6.02	2,620	13.2	5.99	2,880	14.8
PCB-29 PCB-30	5.06		0	5.13 5.13		0	2.94	+	0	3.02 3.02		0	6.02		0	5.99 5.99		0
PCB-31	5.06	2,970	96	5.13	9,100	0 266	2.94 2.94	4,580	0 254	3.02	3,710	213	6.02 6.02	2,280	0 11.48	5.99	2,260	11.61
PCB-34	5.06 5.06	2,970	96	5.13	9,100	0	2.94	4,580	0	3.02	3,710	0	6.02	2,280	0	5.99	2,260	0
PCB-35	5.06	204	7	5.13	242	7	2.94	267	15	3.02	211	12	6.02		0	5.99	244	1.25
PCB-36	5.06		0	5.13	242	0	2.94	207	0	3.02	211	0	6.02		0	5.99	244	0
PCB-37	5.06		1 0	5.13	2,050	60	2.94	1,620	90	3.02	1,070	61	6.02	695	3.50	5.99	861	4.42
PCB-38	5.06		0	5.13	2,030	0	2.94	1,020	0	3.02	1,0,0	0	6.02	033	0	5.99	1 001	0
PCB-39	5.06		0	5.13		0	2.94	1	0	3.02		0	6.02		0	5.99		0
PCB-40	5.06	718	23	5.13	1,030	30	2.94	1,080	60	3.02	771	44	6.02	835	4.21	5.99	769	3.95
PCB-41/64/71/72	5.06		109	5.13	5,090	149	2.94	5,330	295	3.02	3,960	227	6.02	4,210	21.2	5.99	3,810	19.6
PCB-42/59	5.06	1,210	39	5.13	2,380	69	2.94	1,990	110	3.02	1,470	84	6.02	1,350	6.80	5.99	1,260	6.47
PCB-43/49	5.06	2,970	96	5.13	9,130	266	2.94	5,450	302	3.02	4,130	237	6.02	4,070	20.5	5.99	3,640	18.7
PCB-44	5.06	3,890	126	5.13	6,390	186	2.94	5,720	317	3.02	4,390	252	6.02	5,490	27.7	5.99	4,830	24.8
PCB-45	5.06	611	20	5.13	755	22	2.94	767	42	3.02	534	31	6.02	693	3.49	5.99	557	2.86
PCB-46	5.06	303	10	5.13	450	13	2.94	523	29	3.02	416	24	6.02	325	1.64	5.99	301	1.55
PCB-47	5.06		0	5.13	5,580	163	2.94	2,690	149	3.02	2,140	123	6.02		0	5.99		0
PCB-48/75	5.06		22	5.13	1,110	32	2.94	685	38	3.02	523	30	6.02	755	3.80	5.99	694	3.57
PCB-50	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-51	5.06		0	5.13	522	15	2.94	560	31	3.02	436	25	6.02	316	1.59	5.99	244	1.25
PCB-52/69	5.06	4,780	154	5.13	8,660	253	2.94	6,570	364	3.02	5,220	299	6.02	8,120	40.9	5.99	7,500	38.5
PCB-53	5.06		19	5.13	966	28	2.94	1,170	65	3.02	819	47	6.02	736	3.71	5.99	658	3.38
PCB-54	5.06		0	5.13	ļ	0	2.94	1	0	3.02	1	0	6.02	ļ	0	5.99	<u> </u>	0
PCB-55	5.06	+	0	5.13	103	3	2.94	 	0	3.02	 	0	6.02	 	0	5.99	 	0
PCB-56/60	5.06	2,400	78	5.13	3,320	97	2.94	4,400	244	3.02	2,830	162	6.02	3,180	16.0	5.99	3,160	16.2

Table B-3
HSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

								HSM	Particulate S	Sample Colle	ection							
		E	vent 2 Attemp	t 2 (12 -7 PR1	35)			E	vent 1 Attemp	t 3 (4-30 PR14	16)			Ev	ent 1 Attempt	1 (6-10-13 PR:	106)	
	P	R1CSOCLYHP-0	02B		PR1HPDUP-02	В	P	R1CSOCLYHP-0	1C		PR1HPDUP-01	С	P	R1CSOCLYHP-0	1A		PR1HPDUP-01	A
Wet weight (gram)		13.9			15.6			5.85			5.8			20.4			19.9	
% solids		36.4			32.9		1	50.2			52.0			29.5			30.1	
Compound Identified	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	: Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L
PCB-57	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-58	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-61/70	5.06	4,540	147	5.13	7,700	225	2.94	6,590	365	3.02	5,030	288	6.02	8,380	42.2	5.99	7,940	40.8
PCB-62	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-63	5.06	153	5	5.13	670	20	2.94	330	18	3.02		0	6.02	270	1.36	5.99		0
PCB-65	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-67	5.06	113	4	5.13	240	7	2.94		0	3.02		0	6.02		0	5.99		0
PCB-68	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-73	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-74	5.06	1,450	47	5.13	3,490	102	2.94	2,340	130	3.02	1,720	99	6.02	2,360	11.9	5.99	2,180	11.2
PCB-76/66	5.06	3,020	98	5.13	7,430	217	2.94	6,080	337	3.02	4,020	231	6.02	5,110	25.7	5.99	5,000	25.7
PCB-77	5.06	 	0	5.13		0	2.94	856	47	3.02	563	32	6.02	924	4.65	5.99	1,010	5.19
PCB-78	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-79	5.06	+	0	5.13		0	2.94		0	3.02		0	6.02	251	1.26	5.99	253	1.30
PCB-80	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-81	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-82	5.06	1,170	38	5.13	1,470	43	2.94	1,550	86	3.02	1,210	69	6.02	2,890	14.6	5.99	2,690	13.8
PCB-83	5.06		0	5.13	,	0	2.94		0	3.02		0	6.02		0	5,99		0
PCB-84/92	5.06	+	116	5.13	4,720	138	2.94	4,010	222	3.02	3,420	196	6.02	8,330	42.0	5.99	8,250	42.4
PCB-85/116	5.06	1,400	45	5.13	1,760	51	2.94	1,980	110	3.02	1,410	81	6.02	2,690	13.5	5.99	2,560	13.2
PCB-86	5.06		0	5.13	2,. 33	0	2.94	2,500	0	3.02	2,120	0	6.02		0	5.99	2,000	0
PCB-87/117/125	5.06	3,400	110	5.13	4,290	125	2.94	3,780	209	3.02	3,150	181	6.02	8,010	40.3	5.99	7,820	40.2
PCB-88/91	5.06	1,060	34	5.13	1,510	44	2.94	1,380	76	3.02	1,190	68	6.02	2,330	11.7	5.99	2,190	11.3
PCB-89	5.06	<u> </u>	0	5.13	129	4	2.94	1,500	0	3.02	1,130	0	6.02	2,000	0	5.99	2,130	0
PCB-90/101	5.06	8,320	269	5.13	11,200	327	2.94	8,740	484	3.02	7,520	431	6.02	20,200	102	5,99	20,100	103
PCB-93	5.06		0	5.13	11,200	0	2.94	5,7 10	0	3.02	7,320	0	6.02	20,200	0	5,99	20,100	0
PCB-94	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-95/98/102	5.06	5,790	187	5.13	7,820	228	2.94	6,140	340	3.02	5,440	312	6.02	14,000	70.5	5.99	12,300	63.2
PCB-96	5.06	 	0	5.13	7,020	0	2.94	0,140	0	3.02	3,440	0	6.02	14,000	0	5.99	12,300	0
PCB-97	5.06	2,490	80	5.13	3,250	95	2.94	3,050	169	3.02	2,440	140	6.02	6,330	31.9	5.99	6,100	31.3
PCB-99	5.06	<u> </u>	106	5.13	4,780	140	2.94	4,060	225	3.02	3,330	191	6.02	7,960	40.1	5.99	7.950	40.8
PCB-100	5.06		0	5.13	7,700	0	2.94	7,000	0	3.02	3,330	0	6.02	,,500	0	5.99	,,550	0
PCB-103	5.06	ļ	0	5.13	+	0	2.94	+	0	3.02	 	0	6.02	1	0	5.99		0
PCB-103	5.06		0	5.13	1	0	2.94		0	3.02		0	6.02	1	0	5.99		0
PCB-105	5.06	<u> </u>	108	5.13	4,050	118	2.94	4,080	226	3.02	3,100	178	6.02	8,250	41.6	5.99	8,120	41.7
PCB-105	5.06	 	255	5.13	10,500	306	2.94	9,370	519	3.02	7,530	432	6.02	20,100	101	5.99	21,000	108
PCB-100/118	5.06	<u> </u>	16	5.13	750	22	2.94	748	41	3.02	564	32	6.02	1,100	5.54	5.99	1,020	5.24
PCB-108/112	5.06	+	13	5.13	524	15	2.94	494	27	3.02	406	23	6.02	935	4.71	5.99	893	4.59
PCB-100/112	5.06	+	317	5.13	12,300	359	2.94	11,400	631	3.02	8,940	513	6.02	20,000	101	5.99	19,900	102
PCB-111/115	5.06	 	6	5.13	192	6	2.94	11,400	0	3.02	0,340	0	6.02	20,000	1.44	5.99	314	1.61
PCB-117/115	5.06	+	0	5.13	192	0	2.94	+	0	3.02	1	0	6.02	200	0	5.99	314	0
PCB-113 PCB-114		+	6	5.13	213		2.94	+	0	3.02	-	0	6.02	557	2.81	5.99	459	2.36
PCB-114 PCB-119	5.06		5			6 7			+		-	-				5.99	+	
	5.06	1	1	5.13	240		2.94	1	0	3.02	 	0	6.02	263	1.32		323	1.66
PCB-120	5.06	<u> </u>	0	5.13	 	0	2.94	+	0	3.02	1	0	6.02	 	0	5.99	-	0
PCB-121	5.06	<u> </u>	0	5.13	440	0	2.94	1	0	3.02	1	0	6.02	+	0	5.99		0
PCB-122	5.06	+	0	5.13	110	3	2.94		0	3.02		0	6.02	224	0	5.99	222	0
PCB-123	5.06	148	5	5.13	179	5	2.94		0	3.02		0	6.02	301	1.52	5.99	322	1.65

Table B-3
HSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

	<u> </u>							HSM	Particulate :	Sample Colle	ection							
			vent 2 Attemp	t 2 (12 -7 PR13	35)				vent 1 Attemp					Eve	ent 1 Attempt	1 (6-10-13 PR:	106)	-
	PI	R1CSOCLYHP (PR1HPDUP-02	В	P	R1CSOCLYHP-0)1C	Ī	PR1HPDUP-01	С	P	R1CSOCLYHP-0			PR1HPDUP-01	Ā
Wet weight (gram)		13.9			15.6			5.85			5.8			20.4			19.9	-
% solids		36.4			32.9			50.2			52.0			29.5			30.1	
Compound Identified	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	: Weight gram (dry)	Sample Result	Converted Sample Result pg/L
PCB-124	5.06	379	12	5.13	464	14	2.94	475	26	3.02	364	21	6.02	960	4.84	5.99	969	4.98
PCB-126	5.06		0	5.13		0	2.94		0	3.02		0	6.02	261	1.31	5.99	278	1.43
PCB-127	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-128/162	5.06	1,880	61	5.13	2,320	68	2.94	2,110	117	3.02	1,760	101	6.02	5,210	26.2	5.99	5,050	25.9
PCB-129	5.06	590	19	5.13	741	22	2.94	636	35	3.02	475	27	6.02	1,740	8.76	5.99	1,670	8.58
PCB-130	5.06	666	22	5.13	868	25	2.94	757	42	3.02	584	33	6.02	1,720	8.66	5.99	1,600	8.22
PCB-131	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-132/161	5.06	2,890	93	5.13	3,480	102	2.94	3,090	171	3.02	2,750	158	6.02	7,190	36.2	5.99	7,060	36.3
PCB-133/142	5.06	304	10	5.13	374	11	2.94	309	17	3.02	261	15	6.02	748	3.77	5.99	775	3.98
PCB-134/143	5.06	537	17	5.13	689	20	2.94	611	34	3.02	481	28	6.02	1,480	7.45	5.99	1,540	7.91
PCB-135	5.06	1,180	38	5.13	1,520	44	2.94	1,350	75	3.02	1,310	75	6.02	2,020	10.2	5.99	1,990	10.2
PCB-136	5.06	1,110	36	5.13	1,460	43	2.94	1,180	65	3.02	1,070	61	6.02	1,880	9.47	5.99	1,990	10.2
PCB-137	5.06	460	15	5.13	665	19	2.94	634	35	3.02	406	23	6.02	854	4.30	5.99	1,300	6.68
PCB-138/163/164	5.06	10,100	326	5.13	12,300	359	2.94	11,700	648	3.02	9,580	549	6.02	25,100	126	5.99	24,300	125
PCB-139/149	5.06	6,730	217	5.13	8,730	255	2.94	8,060	446	3.02	7,260	416	6.02	13,700	69.0	5.99	13,200	67.8
PCB-140	5.06		0	5.13		0	2.94		0	3.02	ļ	0	6.02		0.000	5.99		0
PCB-141	5.06	1,870	60	5.13	2,340	68	2.94	2,240	124	3.02	1,950	112	6.02	4,640	23.4	5.99	4,540	23.3
PCB-144	5.06	448	14	5.13	507	15	2.94	477	26	3.02	402	23	6.02	873	4.40	5.99	676	3.47
PCB-145	5.06	1110	0	5.13	4 400	0	2.94	1 2 1 2	0	3.02	1.100	0	6.02	2.500	0	5.99	2.522	0
PCB-146/165	5.06	1,140	37	5.13	1,400	41	2.94	1,240	69	3.02	1,100	63	6.02	2,500	12.6	5.99	2,530	13.0
PCB-147	5.06	170	5	5.13	270	8	2.94	216	0	3.02	<u> </u>	0	6.02	273	1.38	5.99	297	1.53
PCB-148 PCB-150	5.06 5.06		0	5.13 5.13		0	2.94 2.94	1	0	3.02 3.02	1	0	6.02 6.02		0	5.99 5.99		0
PCB-151	5.06	1,850	60	5.13	2,250	66	2.94	2,100	116	3.02	1,930	111	6.02	2,960	14.9	5.99	3,120	16.0
PCB-151	5.06	1,630	0	5.13	2,230	0	2.94	2,100	0	3.02	1,930	0	6.02	2,360	0	5.99	3,120	0
PCB-153	5.06	7,950	257	5.13	9,230	269	2.94	9,110	504	3.02	7.790	447	6.02	16,700	84.1	5.99	18,200	93.5
PCB-154	5.06	7,550	0	5.13	123	4	2.94	3,110	0	3.02	7,750	0	6.02	10,700	0	5.99	10,200	0
PCB-155	5.06		0	5.13	123	0	2.94		0	3.02	1	0	6.02		0	5.99		0
PCB-156	5.06	1,070	35	5.13	1,350	39	2.94	1,250	69	3.02	1.010	58	6.02	3,140	15.8	5.99	3,050	15.7
PCB-157	5.06	269	9	5.13	354	10	2.94	336	19	3.02	271	16	6.02	758	3.82	5.99	711	3.65
PCB-158/160	5.06	1,220	39	5.13	1,520	44	2.94	1,410	78	3.02	1,100	63	6.02	2,950	14.9	5.99	3,050	15.7
PCB-159	5.06		0	5.13	<u> </u>	0	2.94		0	3.02		0	6.02		0	5.99	1	0
PCB-166	5.06		0	5.13	İ	0	2.94		0	3.02	İ	0	6.02	İ	0	5.99	İ	0
PCB-167	5.06	436	14	5.13	537	16	2.94	527	29	3.02	442	25	6.02	1,360	6.85	5.99	1,300	6.68
PCB-168	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-169	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-170	5.06	2,600	84	5.13	2,800	82	2.94	2,900	161	3.02	2,900	166	6.02	5,570	28.1	5.99	5,170	26.6
PCB-171	5.06	658	21	5.13	716	21	2.94	826	46	3.02	677	39	6.02	1,420	7.15	5.99	1,360	6.99
PCB-172	5.06	444	14	5.13	505	15	2.94	589	33	3.02	558	32	6.02	833	4.20	5.99	773	3.97
PCB-173	5.06		0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-174	5.06	2,470	80	5.13	2,680	78	2.94	3,010	167	3.02	2,740	157	6.02	5,140	25.9	5.99	4,970	25.5
PCB-175	5.06	116	4	5.13	137	4	2.94		0	3.02		0	6.02		0	5.99		0
PCB-176	5.06		10	5.13	352	10	2.94	354	20	3.02	308	18	6.02	608	3.06	5.99	547	2.81
PCB-177	5.06	1,500	48	5.13	1,590	46	2.94	1,700	94	3.02	1,670	96	6.02	3,180	16.0	5.99	3,020	15.5
PCB-178	5.06	552	18	5.13	653	19	2.94	719	40	3.02	666	38	6.02	877	4.42	5.99	936	4.81
PCB-179	5.06	1,150	37	5.13	1,250	36	2.94	1,320	73	3.02	1,250	72	6.02	2,030	10.2	5.99	1,920	9.86
PCB-180	5.06	5,600	181	5.13	6,220	182	2.94	6,910	382	3.02	6,430	369	6.02	11,400	57.4	5.99	11,500	59.1

Table B-3
HSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

								HSM	Particulate S	Sample Colle	ection							
		E	vent 2 Attemp	ot 2 (12 -7 PR1:	35)			E	vent 1 Attemp	t 3 (4-30 PR14	16)			Ev	ent 1 Attempt	1 (6-10-13 PR:	106)	
	P	R1CSOCLYHP-C)2B		PR1HPDUP-02	В	PI	R1CSOCLYHP-0	1C		PR1HPDUP 01	С	Pi	R1CSOCLYHP 0	1A		PR1HPDUP-01	A
Wet weight (gram)		13.9			15.6			5.85			5.8			20.4			19.9	
% solids		36.4			32.9			50.2			52.0			29.5			30.1	
Compound Identified	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L	Weight gram (dry)	Sample Result	Converted Sample Result pg/L
PCB-181	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-182/187	5.06	3,410	110	5.13	3,790	111	2.94	4,150	230	3.02	3,730	214	6.02	4,870	24.5	5.99	5,030	25.8
PCB-183	5.06	1,440	47	5.13	1,710	50	2.94	1,890	105	3.02	1,690	97	6.02	2,260	11.4	5.99	2,290	11.8
PCB-184	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-185	5.06	317	10	5.13	333	10	2.94	361	20	3.02	320	18	6.02	519	2.61	5.99	532	2.73
PCB-186	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-188	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-189	5.06	116	4	5.13	118	3	2.94		0	3.02		0	6.02		0	5.99	251	1.29
PCB-190	5.06	468	15	5.13	552	16	2.94	585	32	3.02	492	28	6.02	1,060	5.34	5.99	1,010	5.19
PCB-191	5.06	5	0	5.13	113	3	2.94		0	3.02		0	6.02		0	5.99		0
PCB-192	5.00	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-193	5.06	283	9	5.13	276	8	2.94	309	17	3.02	331	19	6.02	510	2.57	5.99	484	2.49
PCB-194	5.06	,	51	5.13	1,480	43	2.94	1,710	95	3.02	1,430	82	6.02	2,540	12.8	5.99	2,420	12.4
PCB-195	5.06	647	21	5.13	707	21	2.94	667	37	3.02	610	35	6.02	1,310	6.60	5.99	1,050	5.39
PCB-196/203	5.06	1,840	59	5.13	1,820	53	2.94	1,900	105	3.02	1,800	103	6.02	1,900	9.57	5.99	2,080	10.7
PCB-197	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-198	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-199	5.06	1,940	63	5.13	1,750	51	2.94	1,870	104	3.02	1,970	113	6.02	2,060	10.4	5.99	2,110	10.8
PCB-200	5.06	203	7	5.13	242	7	2.94	263	15	3.02	217	12	6.02		0	5.99	292	1.50
PCB-201	5.06	230	7	5.13	227	7	2.94	244	14	3.02	234	13	6.02	353	1.78	5.99	287	1.47
PCB-202	5.06	450	15	5.13	410	12	2.94	414	23	3.02	430	25	6.02	561	2.83	5.99	587	3.02
PCB-204	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-205	5.06	5	0	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-206	5.06	2,250	73	5.13	1,420	41	2.94	1,430	79	3.02	1,210	69	6.02	1,930	9.72	5.99	2,110	10.8
PCB-207	5.06	238	8	5.13		0	2.94		0	3.02		0	6.02		0	5.99		0
PCB-208	5.06	749	24	5.13	441	13	2.94	498	28	3.02	412	24	6.02	608	3.06	5.99	621	3.19
PCB-209	5.06	5	0	5.13		0	2.94	1,130	63	3.02	1,080	62	6.02	1,410	7.10	5.99	1,380	7.09

Table B-4 HSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

								HSM D	issolved S	ample Coll	lection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	138)				nt 1 Attemp					Ever	nt 1 Attempt	1 (6-10-13 P	R107)	
	PR	1CSOCLYHD		_ `	R1HDDUP-0	2B	PR1	1CSOCLYHD	•		R1HDDUP-0	1C	PR1	LCSOCLYHD		PR1HDI		Т
		Sample	Ī		Sample	T	1	Sample	Ī		Sample	T		Sample	T		Sample	
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-1	5.01	18.4	92	4.9	19.3	95	4.92		0	5.02		0	4.86		0	4.9		1
PCB-2	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-3	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1
PCB-4/10	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-5/8	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-6	5.01	25.3	127	4.9	25.7	126	4.92		0	5.02		0	4.86		0	4.9		
PCB-7/9	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-11	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-12/13	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-14	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-15	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-16/32	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-17	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-18	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-19	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-20/21/33	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-22	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-23	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-24/27	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-25	5.01		0	4.9		0	4.92		0	5.02		0	4.86	43.6	212	4.9	44.7	219
PCB-26	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-28	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-29	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-30	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-31	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		C
PCB-34	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-35	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		C
PCB-36	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-37	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-38	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-39	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		C
PCB-40	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		(
PCB-41/64/71/72	5.01		0	4.9		0	4.92		0	5.02		0	4.86	215	1045	4.9	207	1014
PCB-42/59	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-43/49	5.01		0	4.9		0	4.92		0	5.02		0	4.86	195	948	4.9	203	995
PCB-44	5.01		0	4.9		0	4.92		0	5.02		0	4.86	0	0	4.9	251	1230
PCB-45	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-46	5.01	0	0	4.9	10.2	50	4.92		0	5.02	ļ	0	4.86		0	4.9		
PCB-47	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-48/75	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		<u> </u>
PCB-50	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-51	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-52/69	5.01		0	4.9		0	4.92		0	5.02		0	4.86	346	1682	4.9	362	1774
PCB-53	5.01		0	4.9		0	4.92		0	5.02		0	4.86	0	0	4.9	53.7	263
PCB-54	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		
PCB-55	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		<u> </u>
PCB-56/60	5.01	I	0	4.9		0	4.92		0	5.02		0	4.86	129	627	4.9	128	627

Table B-4 HSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

								HSM D	issolved S	ample Coll	lection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	138)					t 3 (4-30 PR				Ever	t 1 Attempt	1 (6-10-13 P	R107)	
	PR1	LCSOCLYHD-	02B	PI	R1HDDUP-0	2B	PR1	.CSOCLYHD-	01C	PI	R1HDDUP-0	1C	PR1	LCSOCLYHD	-01A	PR1HDI	OUP-01A	
	Volume	Sample Result		Volume	Sample Result		Volume	Sample Result		Volume	Sample Result		Volume	Sample Result		Volume	Sample Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-57	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-58	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-61/70	5.01		0	4.9		0	4.92		0	5.02		0	4.86	309	1502	4.9	296	1450
PCB-62	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-63	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-65	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-67	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-68	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-73	5.01		0	4.9		0	4.92		0	5.02		0	4.86	400	0	4.9	00.0	174
PCB-74	5.01		0	4.9		0	4.92		0	5.02		0	4.86	102	496	4.9	96.8	474
PCB-76/66 PCB-77	5.01 5.01		0	4.9 4.9		0	4.92 4.92		0	5.02 5.02		0	4.86 4.86	207	1006	4.9 4.9	213	1044
			0						0								-	+ 0
PCB-78 PCB-79	5.01 5.01		0	4.9		0	4.92 4.92		0	5.02 5.02		0	4.86		0	4.9 4.9		+ 0
PCB-79 PCB-80	5.01		0	4.9 4.9		0	4.92 4.92		0	5.02		0	4.86 4.86		0	4.9	-	+ 0
PCB-81	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-82	5.01		0	4.9		0	4.92	18.4	91	5.02	18.5	93	4.86	58.3	283	4.9	70.6	346
PCB-83	5.01		0	4.9		0	4.92	16.4	0	5.02	16.5	0	4.86	36.3	0	4.9	70.6	340
PCB-84/92	5.01		0	4.9		0	4.92		0	5.02		0	4.86	197	957	4.9	208	1019
PCB-85/116	5.01	25.6	128	4.9	24.1	118	4.92	21.9	108	5.02	22.9	115	4.86	47.6	231	4.9	58.6	287
PCB-86	5.01	25.6	0	4.9	24.1	0	4.92	21.9	0	5.02	22.9	0	4.86	47.0	0	4.9	36.0	207
PCB-87/117/125	5.01		0	4.9		0	4.92	50.6	249	5.02	-	0	4.86	193	938	4.9	195	956
PCB-88/91	5.01		0	4.9		0	4.92	21.9	108	5.02	19.7	99	4.86	54.2	263	4.9	61.1	299
PCB-89	5.01		0	4.9		0	4.92	21.9	0	5.02	19.7	0	4.86	34.2	0	4.9	01.1	299
PCB-90/101	5.01	189	947	4.9	193	946	4.92	0	0	5.02	129	648	4.86	466	2265	4.9	488	2391
PCB-93	5.01	189	0	4.9	193	0	4.92	-	0	5.02	129	048	4.86	400	0	4.9	400	2391
PCB-94	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-95/98/102	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-96	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-97	5.01		0	4.9		0	4.92		0	5.02		0	4.86	156	758	4.9	151	740
PCB-99	5.01	66	331	4.9	66.3	325	4.92	52.6	259	5.02	55.7	280	4.86	177	860	4.9	185	907
PCB-100	5.01	- 33	0	4.9	00.5	0	4.92	32.0	0	5.02	33.7	0	4.86		0	4.9	100	0
PCB-103	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-104	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		1 0
PCB-105	5.01		0	4.9		0	4.92	44.6	219	5.02	43.9	220	4.86	177	860	4.9	182	892
PCB-106/118	5.01	0	0	4.9	144	706	4.92	123	605	5.02	105	527	4.86	401	1949	4.9	405	1985
PCB-107/109	5.01	10.8	54	4.9	10.9	53	4.92		0	5.02		0	4.86	22.3	108	4.9	0	0
PCB-108/112	5.01		0	4.9		0	4.92		0	5.02		0	4.86	22.7	110	4.9	24.9	122
PCB-110	5.01		0	4.9		0	4.92	149	733	5.02	146	733	4.86	423	2056	4.9	457	2239
PCB-111/115	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-113	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-114	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-119	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-120	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-121	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-122	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-123	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0

Table B-4 HSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

								HSM D	issolved S	ample Col	lection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	138)				nt 1 Attemp					Even	t 1 Attempt	1 (6-10-13 P	R107)	
	PR:	LCSOCLYHD-	.	_ `	R1HDDUP-0	2B	PR1	CSOCLYHD-		•	R1HDDUP-0	1C	PR1	LCSOCLYHD-			OUP-01A	
	Volume	Sample Result		Volume	Sample Result		Volume	Sample Result		Volume	Sample Result		Volume	Sample Result		Volume	Sample Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-124	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-126	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-127	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-128/162	5.01	27.8	139	4.9	27.6	135	4.92	21.9	108	5.02	20.7	104	4.86	82.2	399	4.9	81.6	400
PCB-129	5.01	0	0	4.9	10.8	53	4.92		0	5.02		0	4.86	33.6	163	4.9	27.8	136
PCB-130	5.01	10.4	52	4.9	10.9	53	4.92		0	5.02		0	4.86	25.7	125	4.9	0	0
PCB-131	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-132/161	5.01	42.2	211	4.9	47	230	4.92		0	5.02		0	4.86	122	593	4.9	128	627
PCB-133/142	5.01		0	4.9		0	4.92		0	5.02	-	0	4.86	20.0	0	4.9	20.4	0
PCB-134/143	5.01	100	0	4.9		0	4.92	40.4	0	5.02		0	4.86	29.8	145	4.9	32.1	157
PCB-135	5.01	19.9	100	4.9	20.3	99	4.92	19.1	94	5.02	20.7	104	4.86	48.2	234	4.9	43.5	213
PCB-136	5.01	44.7	0	4.9	12.0	0	4.92	21.6	106	5.02	17.6	88	4.86	41.9	204	4.9	40	196
PCB-137	5.01	11.7	59	4.9	12.9	63	4.92	126	0	5.02	444	0	4.86	426	0	4.9	426	0
PCB-138/163/164	5.01	162	812	4.9	166	813	4.92	126	620	5.02	114	572	4.86	426	2070	4.9	426	2087
PCB-139/149	5.01		0	4.9		0	4.92	114	561	5.02	118	592	4.86	300	1458	4.9	249	1220
PCB-140	5.01		0	4.9		0	4.92		0	5.02		0	4.86	03.0	0	4.9	02	407
PCB-141 PCB-144	5.01		0	4.9		0	4.92 4.92		0	5.02 5.02		0	4.86	83.8	407	4.9 4.9	83	407 0
PCB-144 PCB-145	5.01 5.01		0	4.9		0	4.92		0	5.02		0	4.86 4.86		0	4.9		0
	_		0	4.9		<u> </u>			0		-	0		50.2	244	4.9	51.7	253
PCB-146/165 PCB-147	5.01 5.01		0	4.9 4.9		0	4.92 4.92		0	5.02 5.02		0	4.86 4.86	50.2	0	4.9	51.7	255
PCB-147	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-140	5.01		0	4.9		0	4.92		0	5.02	-	0	4.86		0	4.9		0
PCB-151	5.01		0	4.9		0	4.92	0	0	5.02	31.3	157	4.86		0	4.9		0
PCB-152	5.01		0	4.9		0	4.92		0	5.02	31.3	0	4.86		0	4.9		1 0
PCB-153	5.01		0	4.9		0	4.92	108	531	5.02	101	507	4.86	360	1750	4.9	346	1695
PCB-154	5.01		0	4.9		0	4.92	100	0	5.02	101	0	4.86	300	0	4.9	340	1033
PCB-155	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-156	5.01		0	4.9		0	4.92	12.8	63	5.02	10.8	54	4.86	46.7	227	4.9	44.1	216
PCB-157	5.01		0	4.9		0	4.92	12.0	0	5.02	10.0	0	4.86	10.7	0	4.9	1112	0
PCB-158/160	5.01		0	4.9		0	4.92		0	5.02		0	4.86	48.3	235	4.9	53.5	262
PCB-159	5.01		0	4.9		0	4.92		0	5.02		0	4.86	70.0	0	4.9		0
PCB-166	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-167	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-168	5.01		0	4.9		0	4.92		0	5.02	<u> </u>	0	4.86		0	4.9		0
PCB-169	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-170	5.01		0	4.9		0	4.92	31.1	153	5.02	29.4	148	4.86	92	447	4.9	101	495
PCB-171	5.01		0	4.9		0	4.92		0	5.02		0	4.86	23.5	114	4.9	0	0
PCB-172	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-173	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-174	5.01		0	4.9		0	4.92	31.6	155	5.02	32	161	4.86	94.3	458	4.9	102	500
PCB-175	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-176	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-177	5.01		0	4.9		0	4.92	19.1	94	5.02	18.6	93	4.86	55.8	271	4.9	48.9	240
PCB-178	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-179	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-180	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0

Table B-4 HSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

								HSM D	issolved S	ample Col	lection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	138)			Eve	nt 1 Attemp	t 3 (4-30 PR	147)			Even	t 1 Attempt	1 (6-10-13 P	R107)	
	PR1	1CSOCLYHD-	02B	P	R1HDDUP-0	2B	PR1	CSOCLYHD-	01C	P	R1HDDUP-0	1C	PR:	1CSOCLYHD-	-01A	PR1HDE	OUP-01A	
		Sample			Sample			Sample			Sample			Sample			Sample	
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-181	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-182/187	5.01		0	4.9		0	4.92	47.8	235	5.02	44.2	222	4.86		0	4.9		0
PCB-183	5.01		0	4.9		0	4.92	20.1	99	5.02	19.9	100	4.86		0	4.9		0
PCB-184	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-185	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-186	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-188	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-189	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-190	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-191	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-192	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-193	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-194	5.01		0	4.9		0	4.92	14.7	72	5.02	15.3	77	4.86		0	4.9		0
PCB-195	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-196/203	5.01		0	4.9		0	4.92	23	113	5.02	18.3	92	4.86		0	4.9		0
PCB-197	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-198	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-199	5.01		0	4.9		0	4.92	19.4	95	5.02	17.9	90	4.86		0	4.9		0
PCB-200	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-201	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-202	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-204	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-205	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-206	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-207	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-208	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0
PCB-209	5.01		0	4.9		0	4.92		0	5.02		0	4.86		0	4.9		0

Table B-5 LSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

								LSM Pa	articulate S	ample Col	lection							
		Eve	ent 2 Attemp	t 2 (12 -7 PR	140)			Eve	nt 1 Attempt	3 (4-30 PR	R149)			Even	t 1 Attempt 1	. (6-10-13 I	PR109)	
	PR:	1CSOCLYLP-	-02B	P	R1LPDUP-0	2B	PR	1CSOCLYLP-	01C	P	R1LPDUP-0	1C	PR	1CSOCLYLP-	01A	P	R1LPDUP-0	1A
Total liters filtered (L)		4.819			4.864			5.009			5.058			4.957			4.844	
		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample
	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result
Compound Identified	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L
PCB-1	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-2	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-3	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-4/10	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	1,230	45	0.305		0
PCB-5/8	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-6	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-7/9	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-11	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-12/13	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-14	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-15	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-16/32	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	3,680	136	0.305	1,200	76
PCB-17	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	2,070	76	0.305	757	48
PCB-18	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	4,560	168	0.305	1,670	105
PCB-19	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	924	34	0.305	342	22
PCB-20/21/33	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	2,910	107	0.305	872	55
PCB-22	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	2,310	85	0.305	682	43
PCB-23	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-24/27	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-25	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	916	34	0.305		0
PCB-26	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	1,390	51	0.305	462	29
PCB-28	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	6,420	237	0.305	1,990	125
PCB-29	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-30	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-31	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	6,260	231	0.305	1,890	119
PCB-34	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-35	0.041		0	0.0657		0	0.0401	1,540	12	0.0405		0	0.183		0	0.305		0
PCB-36	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-37	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	1,850	68	0.305	556	35
PCB-38	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-39	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-40	0.041		0	0.0657		0	0.0401	7,030	56	0.0405		0	0.183	1,610	59	0.305	572	36
PCB-41/64/71/72	0.041		0	0.0657		0	0.0401	31,700	254	0.0405		0	0.183	8,520	315	0.305	2,680	169
PCB-42/59	0.041		0	0.0657		0	0.0401	11,100	89	0.0405		0	0.183	2,940	109	0.305	906	57
PCB-43/49	0.041		0	0.0657		0	0.0401	34,100	273	0.0405		0	0.183	7,790	288	0.305	2,500	157
PCB-44	0.041		0	0.0657		0	0.0401	34,400	275	0.0405		0	0.183	10,600	391	0.305	3,440	217
PCB-45	0.041		0	0.0657		0	0.0401	5,830	47	0.0405		0	0.183	1,290	48	0.305	386	24
PCB-46	0.041		0	0.0657		0	0.0401	3,550	28	0.0405		0	0.183	662	24	0.305		0
PCB-47	0.041		0	0.0657		0	0.0401	14,400	115	0.0405		0	0.183	2,410	89	0.305		0
PCB-48/75	0.041		0	0.0657		0	0.0401	6,340	51	0.0405		0	0.183	1,500	55	0.305	487	31
PCB-50	0.041		0	0.0657		0	0.0401	1,300	10	0.0405		0	0.183		0	0.305		0
PCB-51	0.041		0	0.0657		0	0.0401	2,900	23	0.0405		0	0.183	667	25	0.305		0
PCB-52/69	0.041		0	0.0657		0	0.0401	45,200	362	0.0405		0	0.183	15,500	572	0.305	5,110	322
PCB-53	0.041		0	0.0657		0	0.0401	6,630	53	0.0405		0	0.183	1,550	57	0.305	510	32
PCB-54	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0

Table B-5 LSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

								LSM P	articulate S	ample Col	lection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	140)			Eve	nt 1 Attempt	3 (4-30 PR	(149)			Event	1 Attempt 1	(6-10-13 F	PR109)	
	PR:	1CSOCLYLP-	02B	Р	R1LPDUP-0	2B	PR	1CSOCLYLP-	01C	P	R1LPDUP-0	1C	PR	1CSOCLYLP-	01A	P	R1LPDUP-0	1A
Total liters filtered (L)		4.819			4.864			5.009			5.058			4.957			4.844	
		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample
	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result
Compound Identified	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L
PCB-55	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-56/60	0.041		0	0.0657		0	0.0401	27,600	221	0.0405		0	0.183	6,910	255	0.305	2,050	129
PCB-57	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-58	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-61/70	0.041		0	0.0657		0	0.0401	45,500	364	0.0405		0	0.183	15,700	580	0.305	4,930	310
PCB-62	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-63	0.041		0	0.0657		0	0.0401	1,950	16	0.0405		0	0.183		0	0.305		0
PCB-65	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-67	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-68	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-73	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-74	0.041		0	0.0657		0	0.0401	16,800	134	0.0405		0	0.183	4,730	175	0.305	1,500	94
PCB-76/66	0.041		0	0.0657		0	0.0401	35,700	286	0.0405		0	0.183	10,700	395	0.305	3,220	203
PCB-77	0.041		0	0.0657		0	0.0401	4,370	35	0.0405		0	0.183	1,580	58	0.305	467	29
PCB-78	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-79	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-80	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-81	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-82	0.041		0	0.0657		0	0.0401	8,130	65	0.0405	3,340	27	0.183	4,460	165	0.305	1,260	79
PCB-83	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-84/92	0.041		0	0.0657		0	0.0401	23,700	190	0.0405	8,300	66	0.183	12,600	465	0.305	4,170	263
PCB-85/116	0.041		0	0.0657		0	0.0401	9,720	78	0.0405	3,830	31	0.183	4,210	155	0.305	1,360	86
PCB-86	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-87/117/125	0.041		0	0.0657	7,180	97	0.0401	19,800	159	0.0405	8,330	67	0.183	11,500	425	0.305	3,710	234
PCB-88/91	0.041		0	0.0657		0	0.0401	8,370	67	0.0405	3,320	27	0.183	3,680	136	0.305	1,210	76
PCB-89	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-90/101	0.041		0	0.0657		0	0.0401	49,600	397	0.0405	20,400	163	0.183	30,700	1,133	0.305	10,300	649
PCB-93	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-94	0.041		0	0.0657		0	0.0401		0	0.0405	45.000	0	0.183		0	0.305		0
PCB-95/98/102	0.041		0	0.0657		0	0.0401	37,800	303	0.0405	15,200	122	0.183	20,900	772	0.305	7,250	456
PCB-96	0.041		0	0.0657		0	0.0401	45.000	0	0.0405	6 200	0	0.183	0.000	0	0.305	2 222	0
PCB-97	0.041		0	0.0657		0	0.0401	15,900	127	0.0405	6,290	50	0.183	9,390	347	0.305	2,890	182
PCB-99	0.041		0	0.0657		0	0.0401	21,700	174	0.0405	8,040	64	0.183	11,200	413	0.305	3,690	232
PCB-100	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-103	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-104	0.041		0	0.0657	7.470	0	0.0401	10.300	0	0.0405	7.670	0	0.183	11.000	0	0.305	2.420	0
PCB-105 PCB-106/118	0.041 0.041		0	0.0657 0.0657	7,470 16,800	101 227	0.0401 0.0401	18,300 46,900	147 375	0.0405 0.0405	7,670 19,500	61 156	0.183 0.183	11,600 29,000	428 1,071	0.305 0.305	3,430 9,100	216 573
					10,800						· ·			<u> </u>			· ·	
PCB-107/109 PCB-108/112	0.041 0.041		0	0.0657 0.0657		0	0.0401 0.0401	3,600 2,910	29 23	0.0405 0.0405	1,570	13	0.183 0.183	1,750 1,260	65 47	0.305 0.305	592 441	37 28
PCB-108/112 PCB-110	0.041	23,800	202	0.0657		0	0.0401	59,600	477	0.0405	25,500	204	0.183	31,200	1,152	0.305	10,200	642
PCB-111/115	0.041	23,800	0	0.0657		0	0.0401	1,490	12	0.0405	23,300	0	0.183	650	24	0.305	10,200	0
PCB-117/115	0.041		0	0.0657		0	0.0401	1,490	0	0.0405	-	0	0.183	030	0	0.305	-	0
PCB-113	0.041		0	0.0657		0	0.0401	1,400	11	0.0405	-	0	0.183	658	24	0.305	-	0
PCB-114 PCB-119	0.041		0	0.0657		0	0.0401	1,400	0	0.0405	-	0	0.183	1 030	0	0.305	-	0

Table B-5 LSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

								LSM Pa	articulate S	ample Col	lection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	140)			Eve	nt 1 Attempt	3 (4-30 PR	(149)			Event	1 Attempt 1	(6-10-13 F	PR109)	
	PR:	1CSOCLYLP-	02B	Р	R1LPDUP-0	2B	PR	1CSOCLYLP-	01C	P	R1LPDUP-0	1C	PR:	1CSOCLYLP-	01A	P	R1LPDUP-0	1A
Total liters filtered (L)		4.819			4.864			5.009			5.058			4.957			4.844	
		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample		Sample	Converted Sample
	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result
Compound Identified	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L
PCB-120	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-121	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-122	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-123	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	628	23	0.305		0
PCB-124	0.041		0	0.0657		0	0.0401	2,360	19	0.0405		0	0.183	1,450	54	0.305	446	28
PCB-126	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-127	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-128/162	0.041		0	0.0657	4,140	56	0.0401	9,740	78	0.0405	4,220	34	0.183	6,490	240	0.305	1,940	122
PCB-129	0.041		0	0.0657		0	0.0401	3,070	25	0.0405	1,500	12	0.183	2,170	80	0.305	573	36
PCB-130	0.041		0	0.0657		0	0.0401	3,500	28	0.0405	1,620	13	0.183	2,310	85	0.305	665	42
PCB-131	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-132/161	0.041		0	0.0657	6,000	81	0.0401	14,000	112	0.0405	6,780	54	0.183	9,660	357	0.305	3,040	191
PCB-133/142	0.041		0	0.0657		0	0.0401	1,790	14	0.0405		0	0.183	962	36	0.305	352	22
PCB-134/143	0.041		0	0.0657		0	0.0401	2,820	23	0.0405	1,270	10	0.183	1,880	69	0.305	531	33
PCB-135	0.041		0	0.0657		0	0.0401	9,070	73	0.0405	4,160	33	0.183	2,970	110	0.305	966	61
PCB-136	0.041		0	0.0657		0	0.0401	7,700	62	0.0405	3,630	29	0.183	2,890	107	0.305	947	60
PCB-137	0.041		0	0.0657		0	0.0401	3,500	28	0.0405	1,350	11	0.183	1,790	66	0.305	615	39
PCB-138/163/164	0.041		0	0.0657	20,800	281	0.0401	56,500	452	0.0405	25,400	203	0.183	32,800	1,211	0.305	9,860	621
PCB-139/149	0.041		0	0.0657		0	0.0401	51,100	409	0.0405	24,100	193	0.183	19,700	727	0.305	6,470	407
PCB-140	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-141	0.041		0	0.0657		0	0.0401	12,400	99	0.0405	4,990	40	0.183	6,340	234	0.305	1,810	114
PCB-144	0.041		0	0.0657		0	0.0401	3,280	26	0.0405	1,530	12	0.183	1,170	43	0.305	349	22
PCB-145	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	0.510	0	0.305	1 100	0
PCB-146/165	0.041		0	0.0657		0	0.0401	6,530	52	0.0405	2,990	24	0.183	3,510	130	0.305	1,100	69
PCB-147	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-148	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-150	0.041		0	0.0657		0	0.0401	45.500	0	0.0405	6 220	0	0.183	4 450	0	0.305	4.440	0
PCB-151	0.041		0	0.0657		0	0.0401	15,500	124	0.0405	6,320	51	0.183	4,450	164	0.305	1,440	91
PCB-152 PCB-153	0.041 0.041		0	0.0657 0.0657		0	0.0401	F0 400	403	0.0405 0.0405	10.000	0 159	0.183 0.183	24 100	890	0.305 0.305	6.000	440
PCB-153	0.041		0	0.0657		0	0.0401 0.0401	50,400	0	0.0405	19,900	0	0.183	24,100	0	0.305	6,990	0
PCB-155	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	-	0	0.305		0
PCB-156	0.041		0	0.0657		0	0.0401	6,020	48	0.0405	2,580	21	0.183	3,730	138	0.305	1,150	72
PCB-157	0.041		0	0.0657		0	0.0401	1,550	12	0.0405	2,380	0	0.183	885	33	0.305	1,130	0
PCB-158/160	0.041		0	0.0657		0	0.0401	6,810	55	0.0405	3,110	25	0.183	3,830	141	0.305	1,110	70
PCB-159	0.041		0	0.0657		0	0.0401	0,810	0	0.0405	3,110	0	0.183	3,830	0	0.305	1,110	0
PCB-166	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-167	0.041		0	0.0657		0	0.0401	2,430	19	0.0405		0	0.183	1,430	53	0.305	476	30
PCB-168	0.041		0	0.0657		0	0.0401	2,750	0	0.0405		0	0.183	2,750	0	0.305	+ ,,,	0
PCB-169	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-170	0.041		0	0.0657	5,490	74	0.0401	17,600	141	0.0405	7,250	58	0.183	6,300	233	0.305	1,850	116
PCB-171	0.041		0	0.0657	3,130	0	0.0401	4,560	37	0.0405	1,990	16	0.183	1,740	64	0.305	461	29
PCB-172	0.041		0	0.0657		0	0.0401	3,370	27	0.0405	1,420	11	0.183	952	35	0.305	1	0
PCB-173	0.041		0	0.0657		0	0.0401	-,	0	0.0405		0	0.183	<u> </u>	0	0.305		0
PCB-174	0.041		0	0.0657		0	0.0401	18,500	148	0.0405	6,750	54	0.183	6,990	258	0.305	1,840	116

Table B-5 LSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

								LSM Pa	articulate S	ample Col	llection							
		Eve	nt 2 Attempt	: 2 (12 -7 PR	140)			Eve	nt 1 Attempt	3 (4-30 PF	R149)			Event	t 1 Attempt 1	. (6-10-13 I	PR109)	
	PR:	1CSOCLYLP-	-02B	P	R1LPDUP-0	2B	PR	1CSOCLYLP-	01C	Р	R1LPDUP-0:	1C	PR:	1CSOCLYLP-	01A	Р	R1LPDUP-0	1A
Total liters filtered (L)		4.819			4.864			5.009			5.058			4.957			4.844	
			Converted			Converted			Converted			Converted			Converted			Converted
		Sample	Sample		Sample	Sample		Sample	Sample		Sample	Sample		Sample	Sample		Sample	Sample
	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result	Weight	Result	Result
Compound Identified	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L	gram	pg/g	pg/L
PCB-175	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-176	0.041		0	0.0657		0	0.0401	2,560	20	0.0405		0	0.183	731	27	0.305		0
PCB-177	0.041		0	0.0657		0	0.0401	10,200	82	0.0405	4,240	34	0.183	4,110	152	0.305	1,090	69
PCB-178	0.041		0	0.0657		0	0.0401	5,090	41	0.0405	1,930	15	0.183	1,020	38	0.305		0
PCB-179	0.041		0	0.0657		0	0.0401	9,850	79	0.0405		0	0.183	2,450	90	0.305	685	43
PCB-180	0.041		0	0.0657		0	0.0401	42,700	342	0.0405	15,600	125	0.183	15,200	561	0.305	4,320	272
PCB-181	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-182/187	0.041		0	0.0657		0	0.0401	30,800	247	0.0405	11,100	89	0.183	6,190	229	0.305	1,750	110
PCB-183	0.041		0	0.0657		0	0.0401	12,400	99	0.0405	4,570	37	0.183	2,990	110	0.305	828	52
PCB-184	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-185	0.041		0	0.0657		0	0.0401	2,500	20	0.0405		0	0.183	694	26	0.305		0
PCB-186	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-188	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-189	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-190	0.041		0	0.0657		0	0.0401	3,410	27	0.0405	1,430	11	0.183	1,220	45	0.305	378	24
PCB-191	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-192	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-193	0.041		0	0.0657		0	0.0401	1,960	16	0.0405		0	0.183	685	25	0.305		0
PCB-194	0.041		0	0.0657		0	0.0401	11,200	90	0.0405	3,390	27	0.183	3,110	115	0.305	906	57
PCB-195	0.041		0	0.0657		0	0.0401	4,570	37	0.0405		0	0.183	1,160	43	0.305	342	22
PCB-196/203	0.041		0	0.0657		0	0.0401	18,400	147	0.0405	4,910	39	0.183	3,260	120	0.305	825	52
PCB-197	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-198	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-199	0.041		0	0.0657		0	0.0401	18,800	151	0.0405	5,080	41	0.183	2,730	101	0.305	736	46
PCB-200	0.041		0	0.0657		0	0.0401	2,680	21	0.0405		0	0.183		0	0.305		0
PCB-201	0.041		0	0.0657		0	0.0401	2,300	18	0.0405		0	0.183		0	0.305		0
PCB-202	0.041		0	0.0657		0	0.0401	3,900	31	0.0405		0	0.183	813	30	0.305		0
PCB-204	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-205	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-206	0.041		0	0.0657		0	0.0401	8,100	65	0.0405		0	0.183	2,680	99	0.305		0
PCB-207	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183		0	0.305		0
PCB-208	0.041		0	0.0657		0	0.0401	2,590	21	0.0405		0	0.183	1,000	37	0.305		0
PCB-209	0.041		0	0.0657		0	0.0401		0	0.0405		0	0.183	1,510	56	0.305	397	25

Table B-6 LSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

	1							LSM D	issolved S	ample Coll	ection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	141)				nt 1 Attemp					Even	t 1 Attempt	1 (6-10-13 P	R110)	
	PR	1CSOCLYLD-			R1LDDUP-0	2B	PR	1CSOCLYLD			R1LDDUP-0	1C	PR1	LCSOCLYLD-	•	•	R1LDDUP-0	1A
Compound Identified	Volume Liters	Sample Result pg/L	Mass pg	Volume Liters	Sample Result pg/L	Mass pg	Volume Liters	Sample Result pg/L	Mass pg	Volume Liters	Sample Result pg/L	Mass pg	Volume Liters	Sample Result pg/L	Mass pg	Volume Liters	Sample Result pg/L	Mass pg
PCB-1	4.84	13.4	65	4.97	16.7	83	5.19		0	5.11		0	4.99		0	4.86		0
PCB-2	4.84	1017	0	4.97	10.7	0	5.19		0	5.11		1 0	4.99		1 0	4.86		0
PCB-3	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-4/10	4.84		0	4.97		0	5.19	120	623	5.11	129	659	4.99		0	4.86		0
PCB-5/8	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-6	4.84		0	4.97		0	5.19		0	5.11		0	4.99		1 0	4.86		0
PCB-7/9	4.84		0	4.97		0	5.19		0	5.11		0	4.99		1 0	4.86		0
PCB-11	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-12/13	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-14	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-15	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-16/32	4.84		0	4.97		0	5.19		0	5.11		1 0	4.99		1 0	4.86		0
PCB-17	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-18	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-19	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-20/21/33	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-22	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-23	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-24/27	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-25	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-26	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-28	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-29	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-30	4.84		0	4.97		0	5.19		0	5.11		1 0	4.99		0	4.86		0
PCB-31	4.84		0	4.97		0	5.19		0	5.11		0	4.99		1 0	4.86		0
PCB-34	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-35	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-36	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-37	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-38	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-39	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-40	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-41/64/71/72	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-42/59	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-43/49	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-44	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-45	4.84	İ	0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-46	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-47	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-48/75	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-50	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-51	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-52/69	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-53	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0

Table B-6 LSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

								LSM D	issolved S	ample Coll	ection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	141)					t 3 (4-30 PR				Even	t 1 Attempt :	l (6-10-13 P	R110)	
	PR	1CSOCLYLD-	•		R1LDDUP-0	2B	PR:	LCSOCLYLD-			R1LDDUP-0:	1C	PR:	1CSOCLYLD-			R1LDDUP-0	1A
		Sample	 		Sample	<u> </u>		Sample	<u> </u>		Sample	Ī		Sample	<u> </u>		Sample	<u> </u>
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-54	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-55	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-56/60	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-57	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-58	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-61/70	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-62	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-63	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-65	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-67	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-68	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-73	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-74	4.84		0	4.97		0	5.19		0	5.11		0	4.99	44.2	221	4.86	46.3	225
PCB-76/66	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-77	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-78	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-79	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-80	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-81	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-82	4.84		0	4.97		0	5.19	11.5	60	5.11	10.7	55	4.99		0	4.86		0
PCB-83	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-84/92	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-85/116	4.84	10.5	51	4.97	11.4	57	5.19	14.1	73	5.11	13.0	66	4.99		0	4.86		0
PCB-86	4.84		0	4.97	ļ	0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-87/117/125	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-88/91	4.84		0	4.97		0	5.19	13.0	67	5.11	12.5	64	4.99		0	4.86		0
PCB-89	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-90/101	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-93	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-94	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-95/98/102	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-96	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-97	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-99	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-100	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-103	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-104	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-105	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-106/118	4.84		0	4.97	-	0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-107/109	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-108/112	4.84		0	4.97	-	0	5.19		0	5.11		0	4.99	-	0	4.86		0
PCB-110	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-111/115	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-113	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0

Table B-6 LSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

								ISM D	issolved Sa	ample Coll	ection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	141)					t 3 (4-30 PR				Even	t 1 Attempt :	1 (6-10-13 P	R110)	
	PR	1CSOCLYLD-			R1LDDUP-02	2B	PR:	LCSOCLYLD-	•		R1LDDUP-0:	1C	PR:	1CSOCLYLD-		_ `	R1LDDUP-0	1A
		Sample			Sample	<u> </u>		Sample			Sample	Ī		Sample	T		Sample	T
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	'
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-114	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-119	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-120	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-121	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-122	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-123	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-124	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-126	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-127	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-128/162	4.84		0	4.97		0	5.19	13.6	71	5.11	12.6	64	4.99		0	4.86		0
PCB-129	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-130	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-131	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-132/161	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-133/142	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-134/143	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-135	4.84		0	4.97		0	5.19	12.7	66	5.11	13.3	68	4.99		0	4.86		0
PCB-136	4.84		0	4.97		0	5.19	12.2	63	5.11	0	0	4.99		0	4.86		0
PCB-137	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-138/163/164	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-139/149	4.84		0	4.97		0	5.19	76.4	397	5.11	67.6	345	4.99		0	4.86		0
PCB-140	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-141	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-144	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-145	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-146/165	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-147	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-148	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-150	4.84 4.84		0	4.97 4.97		0	5.19	10.6	0	5.11	17.0	91	4.99 4.99		0	4.86 4.86		0
PCB-151 PCB-152				,,,,,		0	5.19	19.6	102	5.11	17.8				0	4.86		
PCB-152 PCB-153	4.84 4.84		0	4.97 4.97			5.19		0	5.11		0	4.99 4.99		0	4.86		0
PCB-153	4.84			4.97 4.97		0	5.19			5.11		0	4.99 4.99		0	4.86		_
PCB-155	4.84		0	4.97		0	5.19 5.19		0	5.11 5.11		0	4.99		0	4.86		0
PCB-156	4.84		0	4.97		0	5.19		0	5.11		0	4.99	-	0	4.86		0
PCB-157	4.84		0	4.97	<u> </u>	0	5.19		0	5.11		0	4.99	 	0	4.86		0
PCB-158/160	4.84		0	4.97	-	0	5.19		0	5.11		0	4.99	-	0	4.86		0
PCB-159/160	4.84		0	4.97		0	5.19		0	5.11		0	4.99 4.99		0	4.86		0
PCB-166	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-167	4.84		0	4.97		0	5.19		0	5.11		0	4.99	-	0	4.86		0
PCB-168	4.84		0	4.97	 	0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-169	4.84		0	4.97	 	0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-170	4.84		0	4.97		0	5.19	20.6	107	5.11	15.6	80	4.99		0	4.86		0
PCB-171	4.84		0	4.97	 	0	5.19	20.0	0	5.11	13.0	0	4.99		0	4.86		0

Table B-6 LSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

								LSM D	issolved S	ample Coll	ection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	141)			Eve	nt 1 Attemp	t 3 (4-30 PR	150)			Event	t 1 Attempt 1	l (6-10-13 P	R110)	,
	PR:	1CSOCLYLD-	02B	P	R1LDDUP-0	2B	PR:	1CSOCLYLD-	01C	P	R1LDDUP-0:	1C	PR1	1CSOCLYLD-	01A	P	R1LDDUP-0	1A
		Sample			Sample			Sample			Sample			Sample			Sample	Т
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-172	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-173	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-174	4.84		0	4.97		0	5.19	21.9	114	5.11	18.4	94	4.99		0	4.86		0
PCB-175	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-176	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-177	4.84		0	4.97		0	5.19	11.2	58	5.11	10.6	54	4.99		0	4.86		0
PCB-178	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-179	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-180	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-181	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-182/187	4.84		0	4.97		0	5.19	29.8	155	5.11	28.8	147	4.99		0	4.86		0
PCB-183	4.84		0	4.97		0	5.19	13.4	70	5.11	12.2	62	4.99		0	4.86		0
PCB-184	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-185	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-186	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-188	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-189	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-190	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-191	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-192	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-193	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-194	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-195	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-196/203	4.84		0	4.97		0	5.19	13.0	67	5.11	13.2	67	4.99		0	4.86		0
PCB-197	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-198	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-199	4.84		0	4.97		0	5.19	12.2	63	5.11	11.5	59	4.99		0	4.86		0
PCB-200	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-201	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-202	4.84		0	4.97		0	5.19	<u> </u>	0	5.11		0	4.99		0	4.86		0
PCB-204	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-205	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-206	4.84		0	4.97		0	5.19	<u> </u>	0	5.11		0	4.99		0	4.86		0
PCB-207	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-208	4.84		0	4.97		0	5.19		0	5.11		0	4.99		0	4.86		0
PCB-209	4.84		0	4.97	1	0	5.19	<u> </u>	0	5.11		0	4.99	1	0	4.86		0

Table B-7 Whole Water Analytical Results Phase I Report Addendum – Additional Data Evaluation

	1							Whol	e Water Sa	mple Colle	ection							
		Fve	nt 2 Attemn	t 2 (12 -7 PR	134)		1		nt 1 Attemp				T	Fver	nt 1 Attempt	1 (6-10-13 P	R105)	
	DD1	.csoclyww			R1WWDUP-0	128	DD1	CSOCLYWW			143) R1WWDUP-0	110	DD1	CSOCLYWW	-	<u> </u>	1WWDUP-0	01.0
	FKI	Sample	I	FF	Sample	72.B	FNI	Sample	I	FF	Sample	T	LV1	Sample	T	FF	Sample	11A
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-1	4.67	14.1	66	4.99	19.9	99	5.03		0	4.87			4.64	26.3	122	4.76	24.6	117
PCB-2	4.67		0	4.99		0	5.03		0	4.87			4.64		0	4.76		0
PCB-3	4.67		0	4.99		0	5.03		0	4.87			4.64		0	4.76		0
PCB-4/10	4.67		0	4.99		0	5.03	135	679	4.87	170	828	4.64	135	626	4.76	103	490
PCB-5/8	4.67		0	4.99		0	5.03		0	4.87		0	4.64	164	761	4.76	104	495
PCB-6	4.67	26.6	124	4.99	27	135	5.03		0	4.87		0	4.64	57.7	268	4.76		0
PCB-7/9	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-11	4.67		0	4.99		0	5.03		0	4.87		0	4.64	422	1958	4.76	280	1333
PCB-12/13	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-14	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-15	4.67		0	4.99		0	5.03		0	4.87		0	4.64	78.6	365	4.76	30.6	146
PCB-16/32	4.67		0	4.99		0	5.03		0	4.87	259	1261	4.64	222	1030	4.76	160	762
PCB-17	4.67		0	4.99		0	5.03	130	654	4.87	226	1101	4.64	121	561	4.76	78.2	372
PCB-18	4.67		0	4.99		0	5.03		0	4.87		0	4.64	296	1373	4.76	180	857
PCB-19	4.67	28.3	132	4.99		0	5.03	53.8	271	4.87	85.9	418	4.64	63.9	296	4.76	49	233
PCB-20/21/33	4.67		0	4.99		0	5.03		0	4.87		0	4.64	192	891	4.76	104	495
PCB-22	4.67		0	4.99		0	5.03		0	4.87		0	4.64	127	589	4.76	81.2	387
PCB-23	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-24/27	4.67		0	4.99		0	5.03		0	4.87	41.6	203	4.64	31.1	144	4.76		0
PCB-25	4.67		0	4.99	24.8	124	5.03	41.4	208	4.87	66.6	324	4.64	52	241	4.76	34.1	162
PCB-26	4.67		0	4.99		0	5.03		0	4.87	70.9	345	4.64	81.1	376	4.76	46.3	220
PCB-28	4.67		0	4.99		0	5.03		0	4.87	344	1675	4.64	370	1717	4.76	217	1033
PCB-29	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-30	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-31	4.67		0	4.99		0	5.03		0	4.87		0	4.64	309	1434	4.76	210	1000
PCB-34	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-35	4.67		0	4.99		0	5.03	11.2	56	4.87	17	83	4.64	27	125	4.76		0
PCB-36	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-37	4.67		0	4.99		0	5.03		0	4.87		0	4.64	110	510	4.76	59.9	285
PCB-38	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-39	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-40	4.67		0	4.99		0	5.03		0	4.87	48.1	234	4.64	94.9	440	4.76	56.1	267
PCB-41/64/71/72	4.67		0	4.99		0	5.03	149	749	4.87	238	1159	4.64	449	2083	4.76	251	1195
PCB-42/59	4.67		0	4.99		0	5.03	62.3	313	4.87	95.9	467	4.64	157	728	4.76	74.8	356
PCB-43/49	4.67		0	4.99		0	5.03	163	820	4.87	279	1359	4.64	415	1926	4.76	224	1066
PCB-44	4.67		0	4.99		0	5.03	179	900	4.87	279	1359	4.64	568	2636	4.76	234	1114
PCB-45	4.67		0	4.99		0	5.03		0	4.87	42.7	208	4.64	79.3	368	4.76	45.5	217
PCB-46	4.67		0	4.99	12.3	61	5.03	20.1	101	4.87	26.6	130	4.64	43.3	201	4.76		0
PCB-47	4.67		0	4.99		0	5.03		0	4.87	137	667	4.64	148	687	4.76	85.7	408
PCB-48/75	4.67	22.3	104	4.99	24.4	122	5.03		0	4.87	46.1	225	4.64	75.1	348	4.76	45.7	218
PCB-50	4.67		0	4.99		0	5.03	14.1	71	4.87	15.3	75	4.64		0	4.76		0
PCB-51	4.67		0	4.99		0	5.03		0	4.87	32.1	156	4.64	35.3	164	4.76		0
PCB-52/69	4.67		0	4.99		0	5.03	228	1147	4.87	362	1763	4.64	822	3814	4.76	459	2185
PCB-53	4.67		0	4.99		0	5.03	43.9	221	4.87	67.8	330	4.64	89.3	414	4.76	46.5	221
PCB-54	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-55	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0

Table B-7 Whole Water Analytical Results Phase I Report Addendum – Additional Data Evaluation

								Whol	e Water Sa	mple Colle	ection							
		Fve	nt 2 Attemn	t 2 (12 -7 PR	134)				nt 1 Attemp				I	Fver	nt 1 Attempt	1 (6-10-13 P	R105)	
	DD1	CSOCLYWW		. ` 	R1WWDUP-0	128	DD1	CSOCLYWW		. `	143) R1WWDUP-0	11.0	DD1/	CSOCLYWW		. ` 	1WWDUP-0)1 A
		Sample	I		Sample	72.B T	FIXI	Sample	ı	F F	Sample	T	LIVI	Sample	T	F 15	Sample	1
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-56/60	4.67		0	4.99		0	5.03		0	4.87	189	920	4.64	340	1578	4.76	188	895
PCB-57	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-58	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-61/70	4.67		0	4.99	172	858	5.03	200	1006	4.87	345	1680	4.64	817	3791	4.76	446	2123
PCB-62	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-63	4.67		0	4.99		0	5.03		0	4.87	15.3	75	4.64	23.1	107	4.76		0
PCB-65	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-67	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-68	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-73	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-74	4.67		0	4.99		0	5.03	61	307	4.87	109	531	4.64	242	1123	4.76	137	652
PCB-76/66	4.67		0	4.99	118	589	5.03	150	755	4.87	259	1261	4.64	552	2561	4.76	302	1438
PCB-77	4.67		0	4.99		0	5.03		0	4.87	35.5	173	4.64	72.3	335	4.76	44.7	213
PCB-78	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-79	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-80	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-81	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-82	4.67	46.1	215	4.99	42	210	5.03	45.6	229	4.87	79.9	389	4.64	228	1058	4.76	107	509
PCB-83	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-84/92	4.67	129	602	4.99	114	569	5.03	129	649	4.87	230	1120	4.64	674	3127	4.76	380	1809
PCB-85/116	4.67	48.9	228	4.99	47.1	235	5.03	47.1	237	4.87	93	453	4.64	215	998	4.76	92.8	442
PCB-86	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-87/117/125	4.67	117	546	4.99	113	564	5.03	121	609	4.87	215	1047	4.64	677	3141	4.76	388	1847
PCB-88/91	4.67	40.6	190	4.99	37	185	5.03	40.3	203	4.87	77.8	379	4.64	209	970	4.76	108	514
PCB-89	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-90/101	4.67	309	1443	4.99	283	1412	5.03	288	1449	4.87	525	2557	4.64	1660	7702	4.76	920	4379
PCB-93	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-94	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-95/98/102	4.67	211	985	4.99	200	998	5.03	221	1112	4.87	390	1899	4.64	1180	5475	4.76	677	3223
PCB-96	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-97	4.67	95.4	446	4.99	86.8	433	5.03	90.7	456	4.87	163	794	4.64	520	2413	4.76	299	1423
PCB-99	4.67	114	532	4.99	112	559	5.03	116	583	4.87	214	1042	4.64	607	2816	4.76	341	1623
PCB-100	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-103	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-104	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-105	4.67	122	570	4.99	104	519	5.03	113	568	4.87	209	1018	4.64	684	3174	4.76	355	1690
PCB-106/118	4.67	269	1256	4.99	266	1327	5.03	266	1338	4.87	503	2450	4.64	1560	7238	4.76	821	3908
PCB-107/109	4.67		0	4.99	16.2	81	5.03	19.6	99	4.87	30.3	148	4.64	74.7	347	4.76	40.7	194
PCB-108/112	4.67	15.8	74	4.99	13.4	67	5.03	15.1	76	4.87	30.2	147	4.64	72.6	337	4.76	41	195
PCB-110	4.67	353	1649	4.99	307	1532	5.03	343	1725	4.87	594	2893	4.64	1670	7749	4.76	859	4089
PCB-111/115	4.67		0	4.99		0	5.03		0	4.87		0	4.64	23	107	4.76		0
PCB-113	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-114	4.67		0	4.99		0	5.03		0	4.87	11.7	57	4.64	34.1	158	4.76		0
PCB-119	4.67		0	4.99		0	5.03		0	4.87		0	4.64	22.4	104	4.76		0
PCB-120	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-121	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0

Table B-7 Whole Water Analytical Results Phase I Report Addendum – Additional Data Evaluation

								Whole	e Water Sa	mple Colle	ection							
		Fve	nt 2 Attemp	t 2 (12 -7 PR	134)				nt 1 Attemp					Fver	t 1 Attempt	1 (6-10-13 P	R105)	
	DR1	.csoclyww			R1WWDUP-0	12R	DR1	CSOCLYWW		. `	R1WWDUP-(110	DR1	CSOCLYWW	_		1WWDUP-0	11Δ
		Sample	I		Sample	I	11/1	Sample	1	1 1	Sample	T	11/1	Sample	T		Sample	1
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-122	4.67		0	4.99		0	5.03		0	4.87		0	4.64	15.5	72	4.76		0
PCB-123	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-124	4.67	14.5	68	4.99	12.7	63	5.03	13.2	66	4.87	28.1	137	4.64	73.7	342	4.76	37.1	177
PCB-126	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-127	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-128/162	4.67	60	280	4.99	55.3	276	5.03	62.5	314	4.87	114	555	4.64	376	1745	4.76	184	876
PCB-129	4.67	20	93	4.99	19.7	98	5.03	23	116	4.87	35.2	171	4.64	129	599	4.76	67.1	319
PCB-130	4.67	20	93	4.99	19.9	99	5.03	22.5	113	4.87	47.4	231	4.64	117	543	4.76	48.8	232
PCB-131	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-132/161	4.67	90.7	424	4.99	85.6	427	5.03	97.6	491	4.87	178	867	4.64	532	2468	4.76	274	1304
PCB-133/142	4.67	11.1	52	4.99		0	5.03	10.1	51	4.87	16	78	4.64	61.6	286	4.76	28.6	136
PCB-134/143	4.67	18.4	86	4.99	17.6	88	5.03	18	91	4.87	33.4	163	4.64	120	557	4.76	64.2	306
PCB-135	4.67	40.1	187	4.99	41	205	5.03	50.1	252	4.87	75.7	369	4.64	156	724	4.76	96.2	458
PCB-136	4.67		0	4.99	34.5	172	5.03	41.7	210	4.87	75.7	369	4.64	169	784	4.76	84.3	401
PCB-137	4.67	17.7	83	4.99	13.7	68	5.03	18	91	4.87	32.1	156	4.64	73.7	342	4.76	37.1	177
PCB-138/163/164	4.67	334	1560	4.99	313	1562	5.03	365	1836	4.87	674	3282	4.64	1990	9234	4.76	922	4389
PCB-139/149	4.67	210	981	4.99	206	1028	5.03	267	1343	4.87	467	2274	4.64	1040	4826	4.76	601	2861
PCB-140	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-141	4.67	59.9	280	4.99	62.9	314	5.03	71.8	361	4.87	151	735	4.64	358	1661	4.76	170	809
PCB-144	4.67		0	4.99	11.9	59	5.03	16.1	81	4.87	34.4	168	4.64	57.2	265	4.76	29.9	142
PCB-145	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-146/165	4.67	38.3	179	4.99	34.6	173	5.03	40.9	206	4.87	77.3	376	4.64	200	928	4.76	102	486
PCB-147	4.67		0	4.99		0	5.03		0	4.87	-	0	4.64	22.8	106	4.76		0
PCB-148	4.67		0	4.99		0	5.03		0	4.87	-	0	4.64		0	4.76		0
PCB-150	4.67		0	4.99		0	5.03	71.6	0	4.87	120	0	4.64	255	0	4.76	420	0
PCB-151 PCB-152	4.67		0	4.99		0	5.03	71.6	360	4.87	138	672	4.64 4.64	255	1183	4.76	138	657
PCB-152	4.67 4.67	265	1238	4.99 4.99	243	0 1213	5.03 5.03	286	0 1439	4.87 4.87	566	0 2756	4.64	1440	0 6682	4.76 4.76	690	3284
PCB-153	4.67	265	0	4.99	243	0	5.03	280	0	4.87	366	0	4.64	1440	0	4.76	690	0
PCB-155	4.67		0	4.99		0	5.03		0	4.87	 	0	4.64		0	4.76		0
PCB-156	4.67	37.4	175	4.99	30.5	152	5.03	39.1	197	4.87	72.1	351	4.64	218	1012	4.76	106	505
PCB-157	4.67	11.7	55	4.99	30.5	0	5.03	39.1	0	4.87	14.9	73	4.64	56.6	263	4.76	22.3	106
PCB-158/160	4.67	39.1	183	4.99	36.4	182	5.03	44.7	225	4.87	74.2	361	4.64	243	1128	4.76	118	562
PCB-159	4.67	33.1	0	4.99	30.4	0	5.03	77.7	0	4.87	74.2	0	4.64	243	0	4.76	110	0
PCB-166	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-167	4.67	14.5	68	4.99	13.8	69	5.03	15.8	79	4.87	31.3	152	4.64	95.1	441	4.76	39.2	187
PCB-168	4.67		0	4.99		0	5.03		0	4.87	 	0	4.64		0	4.76		0
PCB-169	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-170	4.67	72.1	337	4.99	72.1	360	5.03	99.9	502	4.87	231	1125	4.64	365	1694	4.76	162	771
PCB-171	4.67	22.3	104	4.99	20.4	102	5.03	26	131	4.87	61.8	301	4.64	102	473	4.76	48	228
PCB-172	4.67	15.3	71	4.99	12.9	64	5.03	17.1	86	4.87	46.5	226	4.64	61.7	286	4.76	25.5	121
PCB-173	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-174	4.67		0	4.99		0	5.03	104	523	4.87	245	1193	4.64	413	1916	4.76	181	862
PCB-175	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-176	4.67		0	4.99		0	5.03	13.1	66	4.87	26.2	128	4.64	44.8	208	4.76		0
PCB-177	4.67	43.3	202	4.99	41.1	205	5.03	60.8	306	4.87	136	662	4.64	250	1160	4.76	108	514

Table B-7 Whole Water Analytical Results Phase I Report Addendum – Additional Data Evaluation

	T							Whol	e Water Sa	mple Colle	ection							
		Eve	nt 2 Attemp	t 2 (12 -7 PR	134)			Eve	nt 1 Attemp	t 3 (4-30 PR	145)			Even	t 1 Attempt	1 (6-10-13 P	R105)	
	PR1	CSOCLYWW	7-02B	PR	R1WWDUP-0	2B	PR1	CSOCLYWW	-01C	PR	1WWDUP-0	1C	PR1	CSOCLYWW	′-01A	PR	1WWDUP-0)1A
		Sample			Sample		1	Sample	I		Sample			Sample			Sample	$\overline{}$
	Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result		Volume	Result	
Compound Identified	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg	Liters	pg/L	Mass pg
PCB-178	4.67	17.8	83	4.99	17.9	89	5.03		0	4.87	53.6	261	4.64	70.8	329	4.76	25.4	121
PCB-179	4.67		0	4.99		0	5.03	47	236	4.87	97	472	4.64	165	766	4.76	73.8	351
PCB-180	4.67		0	4.99		0	5.03	222	1117	4.87	540	2630	4.64	889	4125	4.76	396	1885
PCB-181	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-182/187	4.67		0	4.99		0	5.03	133	669	4.87	302	1471	4.64	388	1800	4.76	163	776
PCB-183	4.67		0	4.99		0	5.03	60.7	305	4.87	131	638	4.64	177	821	4.76	79.1	377
PCB-184	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-185	4.67		0	4.99		0	5.03	13	65	4.87	32.3	157	4.64	43.8	203	4.76		0
PCB-186	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-188	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-189	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-190	4.67		0	4.99		0	5.03	19.1	96	4.87	47.6	232	4.64	99	459	4.76	37.7	179
PCB-191	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-192	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-193	4.67		0	4.99		0	5.03		0	4.87	25.4	124	4.64	66	306	4.76	21	100
PCB-194	4.67		0	4.99		0	5.03	49.2	247	4.87	137	667	4.64	191	886	4.76	80.4	383
PCB-195	4.67	15.8	74	4.99	13.8	69	5.03	21.8	110	4.87	51.9	253	4.64	86.2	400	4.76	29.4	140
PCB-196/203	4.67		0	4.99		0	5.03	54.5	274	4.87	153	745	4.64	152	705	4.76	69.5	331
PCB-197	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-198	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-199	4.67	42	196	4.99	36.6	183	5.03	53	267	4.87	157	765	4.64	165	766	4.76	87	414
PCB-200	4.67		0	4.99		0	5.03		0	4.87	20.1	98	4.64	24.6	114	4.76		0
PCB-201	4.67		0	4.99		0	5.03		0	4.87	22	107	4.64	29.3	136	4.76		0
PCB-202	4.67		0	4.99	11.1	55	5.03	15	75	4.87	36.3	177	4.64	44.8	208	4.76		0
PCB-204	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-205	4.67		0	4.99		0	5.03		0	4.87		0	4.64		0	4.76		0
PCB-206	4.67		0	4.99		0	5.03	35.6	179	4.87	105	511	4.64	132	612	4.76	61	290
PCB-207	4.67		0	4.99		0	5.03		0	4.87	11.2	55	4.64		0	4.76		0
PCB-208	4.67		0	4.99		0	5.03	11.5	58	4.87	30	146	4.64	49	227	4.76		0
PCB-209	4.67		0	4.99		0	5.03		0	4.87		0	4.64	94.7	439	4.76		0

Appendix C

Data Evaluation Summaries and Analytical Results – Aroclor PCBs

Table C⁻1
Summary of Detected PCB Aroclors by Method and Event
Phase I Report Addendum – Additional Data Evaluation

									Percent Incre	ease for HSM	Percent Increase for	Percent Increase for	Percent Increase for
									Compared to C	Other Methods	LSM Compared to	HSM Compared to	HSM Compared to
		Nu	mber of E	etections	(Parent	and Dupli	cate Sam	ple)	for Total Cor	ncentrations	WW for Total	LSM for Particulate	LSM for Dissolved
		Total	Concentr	ations	Partic	ulates			When Detec	ted by Both	Concentrations	Concentrations	Concentrations When
	Event/		(μg/L)		(μg	;/L)	Dissolve	d (µg/L)	Metl	hods	When Detected by	When Detected by	Detected by Both
Analyte	Attempt	HSM	LSM	ww	HSM	LSM	HSM	LSM	LSM	ww	Both Methods	Both Methods	Methods
Aroclor 1254	All	2	0	0	2	0	0	0					
Summary													
9 Aroclors	1/2	1.0	0.0	0.0	1.0	0.0	0.0	0.0					
9 Aroclors	2/2	0.0	0.0	0.0	0.0	0.0	0.0	0.0					
9 Aroclors	Ali	0.5	0.0	0.0	0.5	0.0	0.0	0.00					
Percent of Detected Analytes	Ali	6%	0%	0%	6%	0%	0%	0%					

Conclusions

Samples were analyzed for 9 Aroclor PCBs, however, only compounds that were positively identified during analysis are presented. Positive results were reported for HSM particulate analysis only.

Samples were analyzed for a total of 9 Aroclors, however, only Aroclors that were positively identified during analysis are presented.

Percent increase calcuations not performed since Aroclor PCBs were positively identified only for HSM particulate analysis.

<u>Abbreviations</u>

μg/L = micrograms per liter % = percent HSM = high solids mass LSM = low solids mass

Table C-2
Statistical Comparison of the Number of Detected PCB Aroclors by Method and Event
Phase I Report Addendum – Additional Data Evaluation

		of Detection Concentr	•	Maximum Possible	Fishe	er Exact Test (p-v	alue) ²
Event	HSM	LSM	ww	Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW
1/2	2	0	0	9	0.471	0.471	1.000
2/2	0	0	0	9	1.000	1.000	1.000
All	2	0	0	18	0.459	0.459	1.000

Notes

Conclusion

All methods are similar with respect to the number of detected compounds.

Abbreviations

HSM = high solids mass

LSM = low solids mass

¹ Total number of detections for event includes 2 duplicates and 9 compounds.

² The p-value shown is based on a two-sided Fisher Exact Test. A p-value less than 0.05 is considered evidence of a significant difference among methods compared.

Table C-3 HSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

			HS	M Particulate S	Sample Collection	on		
		Event 1 Attempt	2 (7-1-13 PR116)		!	Event 2 Attempt 2	(12-07-13 PR135)
	PR1CSOC	LYHP-01B	PR1HPD	UP-01B	PR1CSOC	LYHP-02B	PR1HPC	OUP-02B
Wet weight (gram)	30).5	30).5	30	0.0	30	0.0
% Solids	3	5	3	3	4	2	4	1
Compound Identified	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L
Aroclor 1254	130	0.00405	160	0.00469	0	0	0	0
Aroclor 1260	0	0	0	0	0	0	0	0

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>

μg/L = micrograms per liter μg/Kg = micrograms per kilogram

Table C-4
HSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

Compound Identified		HSM Dissolved Sample Collection										
		Event 1 Attempt 2 ((07-01-13 PR	117)	Event 2 Attempt 2 (12-07-13 PR138)							
	PR1CS	OCLYHD-01B	PR1H	IDDUP-01B	PR1CSC	OCLYHD-02B	PR1HDDUP-02B					
	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result				
	Liters	ug/L	Liters	ug/L	Liters	ug/L	Liters	ug/L				
N/A	1.01		0.99		0.95		1.00					

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Table C⁻5 LSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

		LSM Particulate Sample Collection										
		Event 1 Attempt 2 (07-05-13 PR119)					Event 2 Attempt 2 (12-11-13 PR140)					
	PR1CSOC	CLYLP-01B	PR1LPE	OUP-01B	PR1CSOC	CLYLP-02B	PR1LPDUP-02B					
TSS (mg/L)	64	4.8	6	7.1	8	.4	13.5					
Total Liters Filtered (L)	0.9	979	1.0	045	1.0	013	1.038					
Compound Identified	Weight gram	Sample Result eight gram ug/Kg		Sample Result ug/Kg	Weight gram	Sample Result ug/Kg	Weight gram	Sample Result ug/Kg				
N/A	0.0636		0.07		0.0085		0.014					

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a ND result.

<u>Abbreviations</u>

μg/Kg = micrograms per kilogram

Table C-6
LSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

				LSM Dissolved Sa	mple Collec	tion						
		Event 1 Attempt 2 (07-05-13 PR120) Event 2 Attempt 2 (12-11-13 PR141)										
	PR1CS0	OCLYLD-01B	PR1	LDDUP-01B	PR1CSC	DUP-02B						
	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result				
Compound Identified	Liters	ug/L	Liters	Liters ug/L		ug/L	Liters	ug/L				
N/A	1.01	_	1.02		1.01		1.04					

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Table C-7
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

		Whole Water Sample Collection									
		Event 1 Attepmt 2	(07-01-13 PI	R115)	Event 2 Attempt 2 (12-07-13 PR134)						
Compound Identified	PR1CSC	CLYWW-01B	PR1W	/WDUP-01B	PR1CSO	CLYWW-02B	PR1W	WDUP-02B			
	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result			
	Liters	ug/L	Liters	ug/L	Liters	ug/L	Liters	ug/L			
N/A	0.985		0.985		1.05		1.04				

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations



Table D⁻1
Summary of Detected Organochlorine Pesticides Congeners by Method and Event
Phase I Report Addendum – Additional Data Evaluation

Analyte (Pesticides)	Event/ Attempt		er of Det Concenti (pg/L) LSM		Partic	and Dur ulates (/L) LSM	Disso	ample) plved g/L) LSM	Compared to C Total Concer When Det	erence for HSM Other Methods for ntrations (pg/L) ected by Both ethods WW	Percent Difference for LSM Compared to WW for Total Concentrations (pg/L) When Detected by Both Methods	Percent Difference for HSM Compared to LSM for Particulate Concentrations (pg/L) When Detected by Both Methods	Percent Difference for HSM Compared to LSM for Dissolved Concentrations (pg/L) When Detected by Both Methods
HEXACHLOROBENZENE	All	1	0	0	1	0	0	0					
ALPHA BHC (ALPHA HEXACHLOROCYCLOHEXANE)	All	2	3	3	1	0	2	3	-3%	-11%	-6%		-5%
GAMMA BHC (LINDANE)	All	4	4	4	3	2	4	4	5%	2%	-2%	-67%	7%
BETA BHC (BETA HEXACHLOROCYCLOHEXANE)	All	3	2	4	2	1	2	2	0%	-42%	2%	-76%	11%
DELTA BHC (DELTA HEXACHLOROCYCLOHEXANE)	All	0	0	0	0	0	0	0					
HEPTACHLOR	All	3	2	2	1	1	2	1	65%	-12%	-46%		265%
ALDRIN	All	2	2	2	0	1	2	2	-5%	-29%	-25%		56%
OXYCHLORDANE	All	3	3	4	2	3	1	0	-11%	-61%	-48%	-65%	
HEPTACHLOR EPOXIDE	All	4	4	4	4	4	4	4	16%	11%	-3%	-48%	73%
BETA-CHLORDANE	All	4	4	4	4	4	4	4	-6%	13%	21%	-66%	119%
TRANS-NONACHLOR	All	4	4	4	4	4	4	4	4%	11%	6%	-51%	113%
ALPHA-CHLORDANE	All	4	4	4	4	4	4	4	5%	24%	17%	-55%	128%
ALPHA ENDOSULFAN	All	0	0	0	0	0	0	0					
O,P'-DDE	All	0	0	0	0	0	0	0					
P,P'-DDE	All	3	0	0	3	0	0	0					
DIELDRIN	All	4	4	4	4	3	4	4	41%	23%	-12%	-46%	105%
ENDRIN	All	0	0	0	0	0	0	0					
CIS-NONACHLOR	All	4	4	4	4	4	3	3	4%	3%	-4%	-42%	128%
BETA ENDOSULFAN	All	0	0	2	0	0	0	0					
O,P'-DDD	All	0	0	0	0	0	0	0					
O,P'-DDT	All	0	0	0	0	0	0	0					
P,P'-DDD	All	2	0	0	2	0	0	0					
P,P'-DDT	All	0	0	0	0	0	0	0					
ENDOSULFAN SULFATE	All	0	0	0	0	0	0	0					
METHOXYCHLOR	All	2	2	3	0	1	2	2	4%	-25%	-28%		52%
MIREX	All	1	0	0	0	0	1	0					
ENDRIN ALDEHYDE	All	0	0	0	0	0	0	0					
ENDRIN KETONE	All	0	0	0	0	0	0	0					
Summary													
28 Analytes	1/2	15	12	13	10	9.0	12	11	8%	-13%	-14%	-70%	79%
28 Analytes	2/2	11	9.0	12	9.5	7.0	8.0	8.0	8%	5%	3%	-37%	107%
28 Analytes	All	13	11	12	9.8	8.0	9.8	9.3	8%	-5%		-55%	91%
Percent of 28 Detected Analytes	All	45%	38%	43%	35%	29%	35%	33%					

Conclusions

HSM has a higher frequency of detection for both total (45%), particulate (35%), and dissolved (35%) concentrations.

Where detected in both methods, HSM total concentrations are on average 8% greater than total LSM concentrations; however, there is large variability among events.

Where detected in both methods, HSM total concentrations are slightly lower on average (-5%) than WW concentrations; however, there is great variability among events.

Where detected in both methods, LSM total concentrations are slightly lower on average than WW concentrations (-7%).

Where detected in both methods, HSM particulate concentrations are on average lower than LSM particulate concentrations (-55%).

 $Where \ detected \ in \ both \ methods, \ HSM \ dissolved \ concentrations \ are \ on \ average \ 91\% \ greater \ than \ LSM \ dissolved \ concentrations.$

<u>Abbreviation</u>

pg/L = picograms per liter

% = percent

HSM = high solids mass

LSM = low solids mass

Table D-2
Statistical Comparison of the Number of Detected Organochlorine Pesticides by Method and Event Phase I Report Addendum – Additional Data Evaluation

	Wate	r Concentra	ation)	Maximum Possible	Chi-Square Test (p-value) ²				
Event	HSM	LSM	ww	Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW		
1/2	29	24	25	56	0.344	0.449	0.849		
2/2	21	18	23	56	0.552	0.699	0.327		
All	50	42	48	112	0.277	0.788	0.414		

Notes

Conclusion

There is no statistically significant difference among methods with respect to the number of detected analytes.

Abbreviations

HSM = high solids mass

LSM = low solids mass

¹ Total number of detections for event includes 2 duplicates and 28 analytes.

² A p-value less than 0.05 is considered significant and is shaded indicating that the number of detects is significantly different between methods.

Table D-3
HSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

		HSM Particulate Sample Collection											
		Eve	ent 1 Attemp	t 2 (7-1 PR	116)			Ever	nt 2 Attem	pt 2(12-7	PR135)		
	PR:	1CSOCLYH	P-01B	Pl	R1HPDUP-	01B	PR1	CSOCLYHP	-02B	P	R1HPDUP	-02B	
Wet weight (gram)		5.67			5.56			5.67		6.1			
% Solids		51.9			35.8			36.2		32.9			
									Converted				
	Weight	Sample	Converted		Sample	Converted		Sample	Sample		Sample	Converted	
	gram	Result	Sample	Weight	Result	Sample	Weight	Result	Result	Weight	Result	Sample	
Compound Identified	(dry)	pg/g	Result pg/L	gram	pg/g	Result pg/L	gram	pg/g	pg/L	gram	pg/g	Result pg/L	
HEXACHLOROBENZENE	2.94		0	1.99		0	2.05	2670	86	2.01		0	
ALPHA BHC (ALPHA HEXACHLOROCYCLOHEXANE)	2.94		0	1.99		0	2.05	102	3	2.01		0	
GAMMA BHC (LINDANE)	2.94	294	14	1.99	319	10	2.05	342	11	2.01		0	
BETA BHC (BETA HEXACHLOROCYCLOHEXANE)	2.94		0	1.99	268	9	2.05	223	7	2.01		0	
DELTA BHC (DELTA HEXACHLOROCYCLOHEXANE)	2.94		0	1.99		0	2.05		0	2.01		0	
HEPTACHLOR	2.94		0	1.99		0	2.05	680	22	2.01		0	
ALDRIN	2.94		0	1.99		0	2.05		0	2.01		0	
OXYCHLORDANE	2.94		0	1.99	476	15	2.05	554	18	2.01		0	
HEPTACHLOR EPOXIDE	2.94	555	26	1.99	1,690	54	2.05	1,590	51	2.01	1,530	45	
BETA-CHLORDANE	2.94	3,930	181	1.99	10,900	347	2.05	10,000	321	2.01	9,350	273	
TRANS-NONACHLOR	2.94	2,780	128	1.99	7,350	234	2.05	8,080	259	2.01	7,790	228	
ALPHA-CHLORDANE	2.94	5,320	245	1.99	15,200	484	2.05	13,500	433	2.01	13,600	398	
ALPHA ENDOSULFAN	2.94		0	1.99		0	2.05		0	2.01		0	
O,P'-DDE	2.94		0	1.99		0	2.05		0	2.01		0	
P,P'-DDE	2.94	7,840	361	1.99	23,000	732	2.05	21,100	677	2.01		0	
DIELDRIN	2.94	3,680	170	1.99	9,470	301	2.05	5,050	162	2.01	5,550	162	
ENDRIN	2.94		0	1.99		0	2.05		0	2.01		0	
CIS-NONACHLOR	2.94	538	25	1.99	2,750	88	2.05	2,320	74	2.01	2,740	80	
BETA ENDOSULFAN	2.94		0	1.99		0	2.05		0	2.01		0	
O,P'-DDD	2.94		0	1.99		0	2.05		0	2.01		0	
O,P'-DDT	2.94		0	1.99		0	2.05		0	2.01		0	
P,P'-DDD	2.94	29,200	1,346	1.99	102,000	3,246	2.05		0	2.01		0	
P,P'-DDT	2.94		0	1.99		0	2.05		0	2.01		0	
ENDOSULFAN SULFATE	2.94		0	1.99		0	2.05		0	2.01		0	
METHOXYCHLOR	2.94		0	1.99		0	2.05		0	2.01		0	
MIREX	2.94		0	1.99		0	2.05		0	2.01		0	
ENDRIN ALDEHYDE	2.94		0	1.99		0	2.05		0	2.01		0	
ENDRIN KETONE	2.94		0	1.99		0	2.05		0	2.01		0	

Table D-4
HSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

			HSM [Dissolved S	ample Colle	ection		
	Ev	ent 1 Attem	ot 2 (7-1 PR11	L 7)	Eve	nt 2 Attemp	t 2 (12-7 PR1	.38)
	PR1CSOC	LYHD-01B	PR1HDE	OUP-01B	PR1CSOC	LYHD-02B	PR1HDE	UP-02B
		Sample		Sample		Sample		Sample
	Volume	Result	Volume	Result	Volume	Result	Volume	Result
Compound Identified	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L
HEXACHLOROBENZENE	2.49		2.41		2.49		2.42	
ALPHA BHC (ALPHA HEXACHLOROCYCLOHEXANE)	2.49		2.41		2.49	60.3	2.42	63.2
GAMMA BHC (LINDANE)	2.49	291	2.41	290	2.49	153	2.42	150
BETA BHC (BETA HEXACHLOROCYCLOHEXANE)	2.49	131	2.41	128	2.49		2.42	
DELTA BHC (DELTA HEXACHLOROCYCLOHEXANE)	2.49		2.41		2.49		2.42	
HEPTACHLOR	2.49	130	2.41	129	2.49		2.42	
ALDRIN	2.49	65	2.41	56	2.49		2.42	
OXYCHLORDANE	2.49	45	2.41	0	2.49		2.42	
HEPTACHLOR EPOXIDE	2.49	320	2.41	335	2.49	112	2.42	119
BETA-CHLORDANE	2.49	1,870	2.41	1,590	2.49	513	2.42	540
TRANS-NONACHLOR	2.49	774	2.41	935	2.49	311	2.42	320
ALPHA-CHLORDANE	2.49	1,870	2.41	1,830	2.49	591	2.42	622
ALPHA ENDOSULFAN	2.49		2.41		2.49		2.42	
O,P'-DDE	2.49		2.41		2.49		2.42	
P,P'-DDE	2.49		2.41		2.49		2.42	
DIELDRIN	2.49	2,390	2.41	2,290	2.49	480	2.42	456
ENDRIN	2.49		2.41		2.49		2.42	
CIS-NONACHLOR	2.49	252	2.41	0	2.49	80.6	2.42	81.8
BETA ENDOSULFAN	2.49		2.41		2.49		2.42	
O,P¹-DDD	2.49		2.41		2.49		2.42	
O,P¹-DDT	2.49		2.41		2.49		2.42	
P,P'-DDD	2.49		2.41		2.49		2.42	
P,P'-DDT	2.49		2.41		2.49		2.42	
ENDOSULFAN SULFATE	2.49		2.41		2.49		2.42	
METHOXYCHLOR	2.49	380	2.41	375	2.49		2.42	
MIREX	2.49	16.5	2.41	0	2.49		2.42	
ENDRIN ALDEHYDE	2.49		2.41		2.49		2.42	
ENDRIN KETONE	2.49		2.41		2.49		2.42	

Abbreviations

Table D-5
LSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

					LSM P	articulate S	Sample C	Collection	า			
		Eve	ent 1 Attem	ot 2 (7-1 P	R119)			Eve	ent 2 Attempt 2	2 (12-7 PF	R140)	
	PR1	LCSOCLYLI	P-01B	PR1LPDUP-01B 2.550			PF	R1CSOCLY	LP-02B	PI	R1LPDUP-	02B
Total Liters Filtered (L)		2.558						2.43		2.357		
Compound Identified	Weight gram	Sample Result pg/g	Converted Sample Result pg/L	Weight gram	Sample Result pg/g	Converted Sample Result pg/L	Weight gram	Sample Result pg/g	Converted Sample Result pg/L	Weight gram	Sample Result pg/g	Converted Sample Result pg/L
HEXACHLOROBENZENE	0.166	1.0,0	0	0.171	1070	0	0.0204	10,0	0	0.0318	1 10,0	0
ALPHA BHC (ALPHA HEXACHLOROCYCLOHEXANE)	0.166		0	0.171		0	0.0204		0	0.0318		0
GAMMA BHC (LINDANE)	0.166	455	30	0.171	617	41	0.0204		0	0.0318		0
BETA BHC (BETA HEXACHLOROCYCLOHEXANE)	0.166	,,,,,	0	0.171	520	35	0.0204		0	0.0318		0
DELTA BHC (DELTA HEXACHLOROCYCLOHEXANE)	0.166		0	0.171		0	0.0204		0	0.0318		0
HEPTACHLOR	0.166		0	0.171	1,290	87	0.0204		0	0.0318		0
ALDRIN	0.166	772	50	0.171	,	0	0.0204		0	0.0318		0
OXYCHLORDANE	0.166	646	42	0.171		0	0.0204	2,710	23	0.0318	2,110	28
HEPTACHLOR EPOXIDE	0.166	2,600	169	0.171	2,770	186	0.0204	6,060	51	0.0318	4,870	66
BETA-CHLORDANE	0.166	20,200	1,311	0.171	22,100	1,482	0.0204	62,600	526	0.0318	49,800	672
TRANS-NONACHLOR	0.166	8,890	577	0.171	10,800	724	0.0204	39,500	332	0.0318	27,400	370
ALPHA-CHLORDANE	0.166	17,800	1,155	0.171	21,800	1,462	0.0204	67,500	567	0.0318	55,600	750
ALPHA ENDOSULFAN	0.166		0	0.171		0	0.0204		0	0.0318		0
O,P'-DDE	0.166		0	0.171		0	0.0204		0	0.0318		0
P,P'-DDE	0.166		0	0.171		0	0.0204		0	0.0318		0
DIELDRIN	0.166		0	0.171	18,000	1,207	0.0204	27,300	229	0.0318	18,200	246
ENDRIN	0.166		0	0.171		0	0.0204		0	0.0318		0
CIS-NONACHLOR	0.166	1,820	118	0.171	2,480	166	0.0204	11,800	99	0.0318	7,820	106
BETA ENDOSULFAN	0.166		0	0.171		0	0.0204		0	0.0318		0
O,P'-DDD	0.166		0	0.171		0	0.0204		0	0.0318		0
O,P'-DDT	0.166		0	0.171		0	0.0204		0	0.0318		0
P,P'-DDD	0.166		0	0.171		0	0.0204		0	0.0318		0
P,P'-DDT	0.166		0	0.171		0	0.0204		0	0.0318		0
ENDOSULFAN SULFATE	0.166		0	0.171		0	0.0204		0	0.0318		0
METHOXYCHLOR	0.166		0	0.171	3,410	229	0.0204		0	0.0318		0
MIREX	0.166		0	0.171		0	0.0204		0	0.0318		0
ENDRIN ALDEHYDE	0.166		0	0.171		0	0.0204		0	0.0318		0
ENDRIN KETONE	0.166		0	0.171		0	0.0204		0	0.0318		0

Abbreviations

Table D-6
LSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

			LSM D	issolved Sa	ample Colle	ection		
	Eve	ent 1 Attem	ot 2 (7-1 PR1	20)	Eve	nt 2 Attemp	t 2 (12-7 PR1	.41)
	PR1CSOC	LYLD-01B	PR1LDD	UP-01B	PR1CSOC	LYLD-02B	PR1LDD	UP-02B
		Sample		Sample		Sample		Sample
	Volume	Result	Volume	Result	Volume	Result	Volume	Result
Compound Identified	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L
HEXACHLOROBENZENE	2.49		2.5		2.44		2.41	
ALPHA BHC (ALPHA HEXACHLOROCYCLOHEXANE)	2.49	25.8	2.5	0	2.44	66.9	2.41	63.5
GAMMA BHC (LINDANE)	2.49	262	2.5	286	2.44	147	2.41	134
BETA BHC (BETA HEXACHLOROCYCLOHEXANE)	2.49	110	2.5	124	2.44		2.41	
DELTA BHC (DELTA HEXACHLOROCYCLOHEXANE)	2.49		2.5		2.44		2.41	
HEPTACHLOR	2.49	70.9	2.5	0	2.44		2.41	
ALDRIN	2.49	36.8	2.5	40.5	2.44		2.41	
OXYCHLORDANE	2.49		2.5		2.44		2.41	
HEPTACHLOR EPOXIDE	2.49	210	2.5	211	2.44	65	2.41	56.2
BETA-CHLORDANE	2.49	865	2.5	1,020	2.44	210	2.41	204
TRANS-NONACHLOR	2.49	422	2.5	605	2.44	123	2.41	120
ALPHA-CHLORDANE	2.49	1,120	2.5	1,120	2.44	218	2.41	200
ALPHA ENDOSULFAN	2.49		2.5		2.44		2.41	
O,P'-DDE	2.49		2.5		2.44		2.41	
P,P'-DDE	2.49		2.5		2.44		2.41	
DIELDRIN	2.49	1,160	2.5	1,240	2.44	220	2.41	214
ENDRIN	2.49		2.5		2.44		2.41	
CIS-NONACHLOR	2.49	117	2.5	0	2.44	33.6	2.41	33.7
BETA ENDOSULFAN	2.49		2.5		2.44		2.41	
O,P'-DDD	2.49		2.5		2.44		2.41	
O,P'-DDT	2.49		2.5		2.44		2.41	
P,P'-DDD	2.49		2.5		2.44		2.41	
P,P'-DDT	2.49		2.5		2.44		2.41	
ENDOSULFAN SULFATE	2.49		2.5		2.44		2.41	
METHOXYCHLOR	2.49	239	2.5	257	2.44		2.41	
MIREX	2.49		2.5		2.44		2.41	
ENDRIN ALDEHYDE	2.49		2.5		2.44		2.41	
ENDRIN KETONE	2.49		2.5		2.44		2.41	

Abbreviations

Table D-7
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

			Whole	e Water Sa	mple Colle	ection		
	Eve	ent 1 Attemp	ot 2 (7-1 PR1	15)	Eve	nt 2 Attemp	t 2 (12 -7 PR1	L34)
	PR1CSOCL	YWW-01B	PR1WW	DUP-01B	PR1CSOCL	YWW-02B	PR1WW	DUP-02B
		Sample		Sample		Sample		Sample
	Volume	Result	Volume	Result	Volume	Result	Volume	Result
Compound Identified	Liters	pg/L	Liters	pg/L	Liters	pg/L	Liters	pg/L
HEXACHLOROBENZENE	2.45		2.53		2.49		2.42	
ALPHA BHC (ALPHA HEXACHLOROCYCLOHEXANE)	2.45		2.53	26.5	2.49	70.1	2.42	72.7
GAMMA BHC (LINDANE)	2.45	313	2.53	311	2.49	146	2.42	147
BETA BHC (BETA HEXACHLOROCYCLOHEXANE)	2.45	136	2.53	127	2.49	23	2.42	30.6
DELTA BHC (DELTA HEXACHLOROCYCLOHEXANE)	2.45		2.53		2.49		2.42	
HEPTACHLOR	2.45	151	2.53	143	2.49		2.42	
ALDRIN	2.45	82.3	2.53	88.7	2.49		2.42	
OXYCHLORDANE	2.45	46.9	2.53	60.6	2.49	33.4	2.42	44.6
HEPTACHLOR EPOXIDE	2.45	371	2.53	376	2.49	128	2.42	137
BETA-CHLORDANE	2.45	2,020	2.53	1,880	2.49	674	2.42	648
TRANS-NONACHLOR	2.45	1,190	2.53	1,070	2.49	439	2.42	421
ALPHA-CHLORDANE	2.45	2,270	2.53	2,440	2.49	661	2.42	665
ALPHA ENDOSULFAN	2.45		2.53		2.49		2.42	
O,P'-DDE	2.45		2.53		2.49		2.42	
P,P'-DDE	2.45		2.53		2.49		2.42	
DIELDRIN	2.45	2,450	2.53	2,610	2.49	421	2.42	449
ENDRIN	2.45		2.53		2.49		2.42	
CIS-NONACHLOR	2.45	257	2.53	290	2.49	113	2.42	115
BETA ENDOSULFAN	2.45		2.53		2.49	633	2.42	711
O,P'-DDD	2.45		2.53		2.49		2.42	
O,P'-DDT	2.45		2.53		2.49		2.42	
P,P'-DDD	2.45		2.53		2.49		2.42	
P,P'-DDT	2.45		2.53		2.49		2.42	
ENDOSULFAN SULFATE	2.45		2.53		2.49		2.42	
METHOXYCHLOR	2.45	480	2.53	523	2.49		2.42	174
MIREX	2.45		2.53		2.49		2.42	
ENDRIN ALDEHYDE	2.45		2.53		2.49		2.42	
ENDRIN KETONE	2.45		2.53		2.49		2.42	

Abbreviations

Appendix E

Data Evaluation Summaries and Analytical Results – SVOCs

Table E-1
Summary of Detected SVOC Compounds by Method and Event
Phase I Report Addendum – Additional Data Evaluation

	Event/	Total Co	Numbei oncentratic LSM		Ī	and Duplice lites (µg/L)		ed (µg/L) LSM	Percent Increa Compared to Of for Total Cond When Detect Meth LSM	ther Methods centrations ed by Both	Percent Increase for LSM Compared to WW for Total Concentrations When Detected by Both Methods	Percent Increase for HSM Compared to LSM for Particulate Concentrations When Detected by Both Methods	Percent Increase for HSM Compared to LSM for Dissolved Concentrations When Detected by Both Methods
Analyte Acetophenone	Attempt	0	0	0	0	LSIVI 0	0	USIVI 0	LSIVI	ww	Both Wethods	Both Wethods	ivietnoas
Bis(2-ethylhexyl)phthalate	All	4	2	1	3	2	3	0	18%	57%	-19%	-81%	
Butylbenzylphthalate	All	4	1	0	3	0	1	1	129%	31/6	15/0	01/0	65%
Carbazole	All	0	0	0	0	0	0	0	125/0				03/8
Dibenzofuran	All	0	0	0	0	0	0	0	+				
Diethylphthalate	All	3	3	4	0	0	3	3	-4%	-34%	-31%		-4%
	All	2	3	1	2	0	3	1	.,	47%			145%
Di-n-butylphthalate		1	0	0	1	0	0	0	193%	47%	-50%		145%
Di-n-octylphthalate	All	2	0	0	1	0		0	F20/				E40/
4-Methylphenol	All		1	0	1	0	2	1	52%				51% -29%
Phenol	All	1	1		0	U U	1	1	-29%				-29%
Summary												•	
50 Analytes	1/2	5.0	2.5	2.5	2.5	0.0	3.5	2.5	67%	38%			45%
50 Analytes	2/2	3.5	2.0	2.0	2.5	1.0	2.0	1.0	9%	-11%			
50 Analytes	All	4.3	2.3	2.3	2.5	0.5	2.8	1.75	51%	19%	-33%	-81%	37%
Percent of Detected Analytes	All	9%	5%	5%	5%	1%	6%	4%					

Conclusions

Samples were analyzed for a total of 50 SVOC compounds, however, only SVOC compounds that were positively identified during analysis are presented.

HSM has a higher frequency of detection for both total (9%), particulate (5%), and dissolved (6%) concentrations.

 $Where \ detected \ in \ both \ methods, HSM \ total \ concentrations \ are \ on \ average \ 51\% \ greater \ than \ total \ LSM \ concentrations.$

Where detected in both methods, HSM total concentrations are on average 19% greater than WW concentrations.

Where detected in both methods, LSM total concentrations are 33% lower than WW concentrations.

Where detected in both methods, HSM particulate concentrations are on average lower than LSM particulate concentrations (-81%).

Where detected in both methods, HSM dissolved concentrations are on average 37% greater than LSM dissolved concentrations.

Abbreviations

μg/L = micrograms per liter % = percent HSM = high solids mass LSM = low solids mass WW = whole water

Table E-2
Statistical Comparison of the Number of Detected SVOC Compounds by Method and Event
Phase I Report Addendum – Additional Data Evaluation

		of Detection	ons (Total ration)	Maximum Possible		Square Test (p-va	nlue) ²
Event	HSM	LSM	w	Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW
1/2	10	5	5	100	0.180	0.180	1.000
2/2	7	4	4	100	0.352	0.352	1.000
All	17	9	9	200	0.105	0.105	1.000

Notes

Conclusion

There is some evidence that HSM method is better than both other methods with respect to the number of detected compounds, however this apparent difference was not statistically significant at alpha = 0.05.

The LSM and WW methods are similar with respect to the number of detected compounds.

Abbreviations

HSM = high solids mass LSM = low solids mass WW = whole water

¹ Total number of detections for event includes 2 duplicates and 50 compounds.

² A p-value less than 0.05 is considered significant and is shaded indicating that the number of detects is significantly different between methods.

Table E-3
HSM Particulate Analytical Results
Phase I Report Addendum − Additional Data Evaluation

		HSM Particulate Sample Collection											
		Event 1 Attempt	: 2 (7-1-13 PR116)			Event 2 Attempt 2	2 (12-07-13 PR135	5)					
	PR1CSOC	LYHP-01B	PR1HPI	OUP-01B	PR1CSOC	LYHP-02B	PR1HPI	OUP-02B					
Wet weight (gram)	25	9.8	25	9.9	30	0.8	29.7						
% Solids	3	15	3	33	4	2	4	11					
Compound Identified	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L					
4-methylphenol	0	0	4,000	0.117	0	0	0	0					
Di-n-butylphthalate	13,000	0.405	4,200	0.123	0	0	0	0					
Butylbenzylphthalate			37,000	1.09	1,200	0.0447	1,400	0.0509					
Bis(2-ethylhexyl)phthalate			25,000	0.734	12,000	0.447	11,000	0.400					
Dibenzofuran					0	0	0	0					
Diethylphthalate					0	0							
Carbazole					0	0	0	0					
Di-n-octylphthalate					2,000	0.0745							

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>

μg/L = micrograms per liter μg/Kg = micrograms per kilogram

Table E-4
HSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

		HSM Dissolved Sample Collection												
		Event 1 Attempt 2	(07-01-13 PR117	7)		Event 2 Attempt 2	(12-07-13 PR13	8)						
Compound Identified	PR1CSO	CLYHD-01B	PR1HD	DUP-01B	PR1CSO	CLYHD-02B	PR1HDDUP-02B							
	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L						
Phenol	2.42	1.7	2.63	0	2.31	0	2.24	0						
4-Methylphenol	2.42	5.4	2.63	8.6	2.31		2.24							
Di-n-butylphthalate	2.42	2.7	2.63	0	2.31	0	2.24	0						
Butylbenzylphthalate	2.42	2.8	2.63		2.31		2.24							
Bis(2-ethylhexyl)phthalate	2.42	29	2.63		2.31	2.1	2.24	2.3						
Diethylphthalate	2.42		2.63	3.4	2.31	1.3	2.24	1.1						
Acetophenone	2.42		2.63		2.31	0	2.24							

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Table E-5
LSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

					LSM	Particulate S	ample Collec	tion				
		Ev	ent 1 Attempt 2	(07-01-13 PR1	19)			Eve	ent 2 Attempt 2	ppt 2 (12-11-13 PR140)		
	P	R1CSOCLYLP-01	LB		PR1LPDUP-018		Р	R1CSOCLYLP-02	!B	PR1LPDUP-02B		
TSS (mg/L)		64.8 67.1						8.4		13.5		
Total Liters Filtered (L)		2.363		2.420				2.418		2.572		
Compound Identified	Weight gram	Sample Result ug/Kg	Converted Sample Result ug/L	Weight gram	Sample Result ug/Kg	Converted Sample Result ug/L	Weight gram	Sample Result ug/Kg	Converted Sample Result ug/L	Weight gram	Sample Result ug/Kg	Converted Sample Result ug/L
Di-n-butylphthalate	0.154	0	0	0.163 0 0			0.0203			0.0347		
Diethylphthalate	0.154			0.163 0 0			0.0203			0.0347		
Bis(2-ethylhexyl)phthalate	0.154			0.163			0.0203	240,000	2	0.0347	180,000	2.43

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

μg/Kg = micrograms per kilogram mg/L = milligrams per liter

Table E-6
LSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

			LSM Disse	olved Sample C	Collection						
	E	vent 1 Attempt 2	(07-01-13 PR12	0)	Event 2 Attempt 2 (12-11-13 PR141)						
	PR1CSOC	CLYLD-01B	PR1LDI	OUP-01B	PR1CSOC	LYLD-02B	PR1LDDUP-02B				
Compound Identified	Volume L	Sample Result ug/L	Volume L	Sample Result ug/L	Volume L	Sample Result ug/L	Volume L	Sample Result ug/L			
Phenol	2.6	2.4	2.52		2.42	0	2.57	0			
4-Methylphenol	2.6	9.3	2.52		2.42		2.57				
Di-n-butylphthalate	2.6	0	2.52	1.1	2.42	0	2.57	0			
Acetophenone	2.6		2.52	0	2.42	0	2.57	0			
Diethylphthalate	2.6		2.52	3.7	2.42	1.3	2.57	1.1			
Butylbenzylphthalate	2.6		2.52	1.7	2.42		2.57				

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Table E-7
Whole Water Analytical Results
Phase I Report Addendum — Additional Data Evaluation

			1	Nhole Water Sa	mple Collection	า					
		Event 1 Attepmt 2	(07-01-13 PR115)		Event 2 Attempt 2 (12-07-13 PR134)					
	PR1CSOC	LYWW-01B	PR1WW	DUP-01B	PR1CSOCI	YWW-02B	PR1WWDUP-02B				
Compound Identified	Volume L	Sample Result ug/L	Volume L	Sample Result ug/L	Volume L	Sample Result ug/L	Volume L	Sample Result ug/L			
4-Methylphenol	2.37	0	2.5		2.36		2.27				
Diethylphthalate	2.37	3.1	2.5	3.7	2.36	1.3	2.27	1.6			
Di-n-butylphthalate	2.37	2.2	2.5	0	2.36	0	2.27	0			
Bis(2-ethylhexyl)phthalate	2.37	5.3	2.5	8.3	2.36	2.5	2.27	3			
Phenol	2.37		2.5	0	2.36		2.27	0			
Acetophenone	2.37		2.5		2.36	0	2.27				

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>



Data Evaluation Summaries and Analytical Results – SVOC SIM

Table F-1
Summary of Detected SVOC SIM Compounds by Method and Event
Phase I Report Addendum – Additional Data Evaluation

Analyte	Event/ Attempt	Total Cor HSM			,	t and Dupli tes (µg/L) LSM		le) ed (μg/L) LSM	Percent Incre Compared to O for Total Con When Detect Meth LSM	ther Methods centrations ted by Both	Percent Increase for LSM Compared to WW for Total Concentrations When Detected by Both Methods	Percent Increase for HSM Compared to LSM for Particulate Concentrations When Detected by Both Methods	Percent Increase for HSM Compared to LSM for Dissolved Concentrations When Detected by Both Methods
Naphthalene	All	4	4	3	2	0	3	4	-37%	-46%	-17%	Wicthods	-47%
2-Methylnaphthalene	All	4	2	4	4	0	4	2	-23%	-36%	18%		-24%
Acenaphthylene	All	0	2	0	0	0	0	2	25/0	3070	10/0		24/0
Acenaphthene	All	3	4	3	2	0	3	4	-3%	-47%	-1%		-11%
Fluorene	All	4	4	4	3	1	4	4	-20%	-45%	-32%	-94%	35%
Phenanthrene	All	4	2	4	4	2	4	2	-82%	-53%	-37%	-94%	51%
Anthracene	All	3	2	3	3	2	2	2	-84%	-87%	-44%	-95%	-26%
Fluoranthene	All	4	4	4	4	4	3	2	-53%	-71%	-38%	-73%	84%
Pyrene	All	4	4	4	4	4	4	2	-14%	-60%	-42%	-77%	153%
Benzo(a)anthracene	All	4	2	2	4	2	2	1	-90%	-93%	-36%	-95%	191%
Chrysene	All	4	2	2	4	2	2	2	-89%	-93%	-31%	-95%	196%
Benzo(b)fluoranthene	All	4	4	4	4	4	2	1	-75%	-75%	-14%	-78%	301%
Benzo(k)fluoranthene	All	4	4	4	4	4	2	1	-75%	-79%	-21%	-78%	351%
Benzo(a)pyrene	All	4	4	4	4	4	2	0	-71%	-75%	-23%	-74%	
Indeno(1,2,3-cd)pyrene	All	4	3	2	4	3	0	0	-53%	-97%	-43%	-53%	
Dibenzo(a,h)anthracene	All	4	2	1	4	2	0	0	-94%	-97%	-39%	-94%	
Benzo(g,h,i)perylene	All	4	4	3	4	4	0	0	-75%	-55%	28%	-75%	
1-Methylnaphthalene	All	4	4	4	3	0	4	4	-9%	-29%	-16%		-11%
Benzo[e]pyrene	All	4	4	4	4	4	2	1	-74%	-75%	-17%	-76%	239%
Perylene	All	4	2	2	4	2	0	0	-95%	-97%	-44%	-95%	
3,6-Dimethylphenanthrene	All	3	2	1	3	2	2	0	46%	-91%	-97%	-76%	
1-Methylanthracene	All	4	4	4	4	4	4	4	-54%	-34%	21%	-92%	116%
1-Methylfluoranthene	All	4	4	2	4	4	2	1	-83%	-90%	-26%	-87%	101%
1-Methylpyrene	All	3	2	1	3	2	0	0	-95%	-98%	-60%	-95%	
2,6-Dimethylnaphthalene	All	4	4	4	4	3	4	4	12%	-24%	-30%	-90%	77%
2,3,5-Trimethylnaphthalene	All	4	4	4	4	1	4	4	16%	-42%	-45%	-94%	30%
1,1'-Biphenyl	All	1	1	1	0	0	1	1		-14%			
Dibenzofuran	All	2	2	1	2	0	2	2	140%	-76%	-90%		113%
1-Methylphenanthrene	All	4	4	4	3	1.0	4	4	18%	-36%	-33%	-97%	258%
Dibenzothiophene	All	4	2.0	4	2.0	0.0	4.0	2.0	38%	-41%	-82%		31%
Summary													
30 Analytes	1/2	25.5	18	18	21.5	11.5	12.5	9	-26%	-28%	6%	-65%	7%
30 Analytes	2/2	28	28	25.5	27.5	19	22.5	19	-44%	-83%	-47%	-93%	130%
30 Analytes	All	26.75	23	21.75	24.5	15.25	17.5	14	-37%	-60%	-27%	-83%	92%
Percent of Detected Analytes	All	89%	77%	73%	82%	51%	58%	47%					

Conclusions

HSM has a higher frequency of detection for both total (89%), particulate (82%), and dissolved (58%) concentrations.

Where detected in both methods, HSM total concentrations are 37% lower than total LSM concentrations although there is large variability among events

Where detected in both methods, HSM total concentrations are 60% lower than WW concentrations however there is great variability among events

Where detected in both methods, LSM total concentrations are 27% lower than WW concentrations.

Where detected in both methods, HSM particulate concentrations are on average lower than LSM particulate concentrations (-83%).

Where detected in both methods, HSM dissolved concentrations are on average 92% greater than LSM dissolved concentrations.

<u>Abbreviations</u>

μg/L = micrograms per liter % = percent HSM = high solids mass

I ISM - mgn sonds ma

LSM = low solids mass WW = whole water

Table F-2
Statistical Comparison of the Number of Detected SVOC SIM Compounds by Method and Event Phase I Report Addendum – Additional Data Evaluation

	Number o	of Detections (To Concentration)	otal Water	Maximum Possible	Chi-S	Square Test (p-va	alue) ²
Event	HSM	LSM	ww	Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW
1/2	51	36	36	60	0.002	0.002	1.000
2/2	56	56	51	60	1.000	0.001	0.142
All	107	92	87	120	0.010	0.001	0.459

Notes

Conclusion

The HSM method is better than both other methods with respect to the number of detected compounds.

The LSM and WW methods are similar with respect to the number of detected compounds.

Abbreviations

HSM = high solids mass

LSM = low solids mass

WW = whole water

¹ Total number of detections for event includes 2 duplicates and 30 compounds.

² A p-value less than 0.05 is considered significant and is shaded indicating that the number of detects is significantly different between methods.

Table F-3 HSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

			HS	M Particulate	Sample Collect	ion			
		Event 1 Attempt	: 2 (7-1-13 PR116))	E	Event 2 Attempt	2 (12-7-13 PR135	5)	
	PR1CSOC	LYHP-01B	PR1HPE	OUP-01B	PR1CSOC	LYHP-02B	PR1HPE	OUP-02B	
Wet weight (gram)	30).5	30	0.5	30	0.1	30.1		
% Solids		15		33	4		41		
Compound Identified	Sample Result ug/kg	Converted Sample Result ug/L	Sample Result ug/kg	Converted Sample Result ug/L	Sample Result ug/kg	Converted Sample Result ug/L	Sample Result ug/kg	Converted Sample Result ug/L	
Naphthalene					90	0.00335	410	0.0149	
2-Methylnaphthalene	110	0.00342	71	0.002083377	76	0.00283	73	0.00266	
Acenaphthylene									
Acenaphthene					52	0.00194	40	0.00146	
Fluorene	75	0.00233			80	0.00298	66	0.00240	
Phenanthrene	710	0.0221	300	0.00880	790	0.0294	590	0.0215	
Anthracene	120	0.00373			100	0.00373	82	0.0030	
Fluoranthene	1,900	0.0591	770	0.0226	1,000	0.0373	1100	0.0400	
Pyrene	1,000	0.0311	680	0.0200	940	0.0350	810	0.0295	
Benzo(a)anthracene	780	0.0243	310	0.00910	580	0.0216	470	0.0171	
Chrysene	920	0.0286	410	0.0120	940	0.0350	770	0.0280	
Benzo(b)fluoranthene	890	0.0277	390	0.0114	830	0.0309	720	0.0262	
Benzo(k)fluoranthene	730	0.0227	290	0.00851	750	0.0280	630	0.0229	
Benzo(a)pyrene	750	0.0233	280	0.00822	560	0.0209	470	0.0171	
Indeno(1,2,3-cd)pyrene	400	0.0124	180	0.00528	540	0.0201	420	0.0153	
Dibenzo (a, h) anthracene	120	0.00373	66	0.00194	200	0.00745	150	0.00546	
Benzo(g,h,i)perylene	410	0.0128	220	0.00646	650	0.0242	540	0.0196	
1-Methylnaphthalene	68	0.00212			54	0.00201	49	0.00178	
Benzo[e]pyrene	640	0.0199	270	0.00792	650	0.0242	570	0.0207	
Perylene	200	0.00622	77	0.00226	170	0.00634	140	0.00509	
3,6-Dimethylphenanthrene	54	0.00168			53	0.00198	37	0.00135	
1-Methylanthracene	260	0.00809	91	0.00267	110	0.00410	80	0.00291	
1-Methylfluoranthene	180	0.00560	110	0.00323	260	0.00969	210	0.00764	
1-Methylpyrene	87	0.00271			74	0.00276	64	0.00233	
2,6-Dimethylnaphthalene	150	0.00467	100	0.00293	70	0.00261	77	0.00280	
2,3,5-Trimethylnaphthalene	120	0.00373	76	0.00223	53	0.00198	60	0.00218	
Dibenzofuran					48	0.00179	37	0.00135	
1-Methylphenanthrene	190	0.00591			94	0.00350	120	0.00437	
Dibenzothiophene	51	0.00159			52	0.00194	0	0	

A "O" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>

μg/L = micrograms per liter μg/Kg = micrograms per kilogram

Table F-4 HSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

			HSI	/I Dissolved Sa	ample Co	ollection		
	Εν	ent 1 Attempt 2	(07-01-1	3 PR117)	Ev	ent 2 Attempt 2	2 (12-07-1	3 PR138)
	PR1CS	OCLYHD-01B	PR1	HDDUP-01B	PR1CS	OCLYHD-02B	PR1H	IDDUP-02B
Compound Identified	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L
Naphthalene	2.61	0.24	2.52	0.23	2.26	0.035	2.28	
2-Methylnaphthalene	2.61	0.34	2.52	0.31	2.26	0.052	2.28	0.049
Acenaphthylene	2.61		2.52		2.26	0	2.28	0
Acenaphthene	2.61	0.019	2.52		2.26	0.015	2.28	0.013
Fluorene	2.61	0.025	2.52	0.020	2.26	0.030	2.28	0.028
Phenanthrene	2.61	0.076	2.52	0.063	2.26	0.064	2.28	0.060
Anthracene	2.61		2.52		2.26	0.011	2.28	0.0089
Fluoranthene	2.61	0.054	2.52		2.26	0.069	2.28	0.060
Pyrene	2.61	0.083	2.52	0.069	2.26	0.056	2.28	0.058
Benzo(a)anthracene	2.61		2.52		2.26	0.023	2.28	0.020
Chrysene	2.61		2.52		2.26	0.034	2.28	0.032
Benzo(b)fluoranthene	2.61		2.52		2.26	0.033	2.28	0.032
Benzo(k)fluoranthene	2.61		2.52		2.26	0.029	2.28	0.026
Benzo(a) pyrene	2.61		2.52		2.26	0.020	2.28	0.018
1-Methylnaphthalene	2.61	0.23	2.52	0.21	2.26	0.053	2.28	0.047
Benzo[e]pyrene	2.61		2.52		2.26	0.021	2.28	0.019
Perylene	2.61		2.52		2.26	0	2.28	0
3,6-Dimethylphenanthrene	2.61		2.52		2.26	0.011	2.28	0.0095
1-Methylanthracene	2.61	0.050	2.52	0.043	2.26	0.022	2.28	0.016
1-Methylfluoranthene	2.61		2.52		2.26	0.016	2.28	0.013
1-Methylpyrene	2.61		2.52		2.26	0	2.28	0
2,6-Dimethylnaphthalene	2.61	0.14	2.52	0.12	2.26	0.092	2.28	0.087
2,3,5-Trimethylnaphthalene	2.61	0.070	2.52	0.074	2.26	0.052	2.28	0.011
1,1'-Biphenyl	2.61	0.019	2.52		2.26		2.28	
Dibenzofuran	2.61		2.52		2.26	0.016	2.28	0.0094
1-Methylphenanthrene	2.61	0.069	2.52	0.061	2.26	0.036	2.28	0.032
Dibenzothiophene	2.61	0.026	2.52	0.025	2.26	0.018	2.28	0.016

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>

Table F-5
LSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

					LSM	1 Particulate S	ample Collec	tion				
		E	vent 1 Attempt 2	(07-05-13 PR1	19)			Ev	vent 2 Attempt 2	(12-11-13 PR1	40)	
	P	R1CSOCLYLP-0	1B		PR1LPDUP-01E	1	P	R1CSOCLYLP-02	2B	-	PR1LPDUP-02B	}
TSS (mg/L)		64.8			67.1			8.4			13.5	
Total Liters Filtered (L)		2.53			2.46			2.396			2.502	
, ,			Converted			Converted			Converted			Converted
Compound Identified	Weight	Sample Result	Sample Result	Weight	Sample Result	Sample Result	Weight	Sample Result	Sample Result	Weight	Sample Result	Sample Result
	gram	ug/Kg	ug/L	gram	ug/Kg	ug/L	gram	ug/Kg	ug/L	gram	ug/Kg	ug/L
Naphthalene	0.164	0	0	0.165	0	0	0.0204			0.0338		
2-Methylnaphthalene	0.164			0.165	0	0	0.0204			0.0338		
Acenaphthylene	0.164			0.165			0.0204	0	0	0.0338	0	0
Fluorene	0.164			0.165			0.0204			0.0338	6,900	0.0932
Phenanthrene	0.164			0.165			0.0204	2,500	0.0210	0.0338	65,000	0.878
Anthracene	0.164			0.165			0.0204	870	0.00731	0.0338	10,000	0.135
Fluoranthene	0.164	870	0.0564	0.165	1,600	0.107	0.0204	9,100	0.0764	0.0338	130,000	1.76
Pyrene	0.164	930	0.0603	0.165	1,000	0.0671	0.0204	8,400	0.0706	0.0338	91,000	1.23
Benzo(a) anthracene	0.164			0.165			0.0204	6,700	0.0563	0.0338	54,000	0.729
Chrysene	0.164			0.165			0.0204	8,600	0.0722	0.0338	83,000	1.12
Benzo(b) fluoranthene	0.164	630	0.0408	0.165	880	0.0590	0.0204	7,200	0.0605	0.0338	82,000	1.11
Benzo(k) fluoranthene	0.164	500	0.0324	0.165	720	0.0483	0.0204	8,500	0.0714	0.0338	64,000	0.864
Benzo(a) pyrene	0.164	450	0.0292	0.165	540	0.0362	0.0204	6,600	0.0554	0.0338	56,000	0.756
Indeno(1,2,3-cd)pyrene	0.164			0.165	300	0.0201	0.0204	5,100	0.0428	0.0338	44,000	0.594
Dibenzo(a,h) anthracene	0.164			0.165			0.0204	1,800	0.0151	0.0338	16,000	0.216
Benzo(g,h,i)perylene	0.164	310	0.0201	0.165	340	0.0228	0.0204	6,200	0.0521	0.0338	55,000	0.743
Benzo[e]pyrene	0.164	420	0.0272	0.165	550	0.0369	0.0204	7,300	0.0613	0.0338	61,000	0.824
Perylene	0.164			0.165			0.0204	2,000	0.0168	0.0338	15,000	0.203
3,6-Dimethylphenanthrene	0.164			0.165	330	0.0221	0.0204	500	0.0042	0.0338	0	0
1-Methylanthracene	0.164	620	0.0402	0.165	630	0.0423	0.0204	1,700	0.0143	0.0338	15,000	0.203
1-Methylfluoranthene	0.164	310	0.0201	0.165	320	0.0215	0.0204	2,700	0.0227	0.0338	24,000	0.324
1-Methylpyrene	0.164			0.165			0.0204	840	0.00706	0.0338	7,100	0.0959
2,6-Dimethylnaphthalene	0.164	480	0.0311	0.165	450	0.0302	0.0204			0.0338	5,400	0.073
2,3,5-Trimethylnaphthalene	0.164	0	0	0.165	700	0.0470	0.0204	0	0	0.0338	0	0
1-Methylphenanthrene	0.164			0.165			0.0204	0	0	0.0338	10,000	0.135
Dibenzothiophene	0.164			0.165			0.0204			0.0338	0	0

Note

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

µg/Kg = micrograms per kilogram µg/L = micrograms per liter mg/L = milligrams per liter

Table F-6
LSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

			LSN	1 Dissolved Sa	mple Colle	ction		
	Ev	ent 1 Attempt 2	(07-05-13 PR	(120)	Ev	ent 2 Attempt 2	(12-11-13 PF	R141)
	PR1CSO	CLYLD-01B	PR1LD	DUP-01B	PR1CSO	CLYLD-02B	PR1LD	DUP-02B
	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result
Compound Identified	Liters	ug/L	Liters	ug/L	Liters	ug/L	Liters	ug/L
Naphthalene	2.46	0.34	2.46	0.37	2.43	0.051	2.5	0.037
2-Methylnaphthalene	2.46	0.41	2.46	0.44	2.43		2.5	
Acenaphthylene	2.46		2.46		2.43	0.0058	2.5	0.018
Acenaphthene	2.46	0.022	2.46	0.020	2.43	0.014	2.5	0.0072
Fluorene	2.46	0.021	2.46	0.022	2.43	0.021	2.5	0.014
Phenanthrene	2.46		2.46		2.43	0.038	2.5	0.044
Anthracene	2.46		2.46		2.43	0.015	2.5	0.012
Fluoranthene	2.46		2.46		2.43	0.039	2.5	0.031
Pyrene	2.46		2.46		2.43	0.026	2.5	0.019
Benzo(a)anthracene	2.46		2.46		2.43	0.0074	2.5	0
Chrysene	2.46		2.46		2.43	0.014	2.5	0.0083
Benzo(b)fluoranthene	2.46		2.46		2.43	0.0081	2.5	0
Benzo(k)fluoranthene	2.46		2.46		2.43	0.0061	2.5	0
Benzo(a)pyrene	2.46		2.46		2.43	0	2.5	0
Indeno(1,2,3-cd)pyrene	2.46		2.46		2.43	0	2.5	0
Dibenzo(a,h)anthracene	2.46		2.46		2.43	0	2.5	
Benzo(g,h,i)perylene	2.46		2.46		2.43	0	2.5	0
1-Methylnaphthalene	2.46	0.28	2.46	0.31	2.43	0.063	2.5	0.034
Benzo[e]pyrene	2.46		2.46		2.43	0.0059	2.5	0
Perylene	2.46		2.46		2.43	0	2.5	0
3,6-Dimethylphenanthrene	2.46		2.46		2.43	0	2.5	0
1-Methylanthracene	2.46	0.031	2.46	0.030	2.43	0.0087	2.5	0.0049
1-Methylfluoranthene	2.46		2.46		2.43	0.0072	2.5	0
1-Methylpyrene	2.46		2.46		2.43	0	2.5	0
2,6-Dimethylnaphthalene	2.46	0.10	2.46	0.10	2.43	0.053	2.5	0.027
2,3,5-Trimethylnaphthalene	2.46	0.054	2.46	0.054	2.43	0.036	2.5	0.014
1,1'-Biphenyl	2.46		2.46		2.43		2.5	0.0049
Dibenzofuran	2.46		2.46	† †	2.43	0.0073	2.5	0.0046
1-Methylphenanthrene	2.46	0.037	2.46	0.037	2.43	0.0069	2.5	0.0057
Dibenzothiophene	2.46		2.46	† †	2.43	0.011	2.5	0.015

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect ND result.

<u>Abbreviations</u>

Table F-7
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

			W	ole Water Sar	nple Colle	ction		
	Ev	ent 1 Attepmt 2	(07-01-13 P	R115)	Eve	ent 2 Attempt 2	(12-07-134 P	R134)
	PR1CSOC	CLYWW-01B	PR1WV	VDUP-01B	PR1CSOC	CLYWW-01B	PR1WV	VDUP-01B
Compound Identified	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L
Naphthalene	2.49	0.26	2.42	0.30	2.30		2.26	0.23
2-Methylnaphthalene	2.49	0.32	2.42	0.40	2.30	0.044	2.26	0.25
Acenaphthylene	2.49		2.42		2.30	0	2.26	0
Acenaphthene	2.49	0.023	2.42		2.30	0.013	2.26	0.12
Fluorene	2.49	0.031	2.42	0.028	2.30	0.026	2.26	0.18
Phenanthrene	2.49	0.11	2.42	0.097	2.30	0.065	2.26	1.5
Anthracene	2.49	0.022	2.42		2.30	0.013	2.26	0.29
Fluoranthene	2.49	0.15	2.42	0.12	2.30	0.082	2.26	2.9
Pyrene	2.49	0.15	2.42	0.14	2.30	0.066	2.26	1.8
Benzo(a)anthracene	2.49		2.42		2.30	0.032	2.26	1.2
Chrysene	2.49		2.42		2.30	0.050	2.26	1.7
Benzo(b)fluoranthene	2.49	0.05	2.42	0.042	2.30	0.047	2.26	1.8
Benzo(k)fluoranthene	2.49	0.049	2.42	0.043	2.30	0.039	2.26	1.3
Benzo(a)pyrene	2.49	0.038	2.42	0.033	2.30	0.030	2.26	1.3
Indeno(1,2,3-cd)pyrene	2.49		2.42		2.30	0.012	2.26	1.1
Dibenzo(a,h)anthracene	2.49		2.42		2.30		2.26	0.38
Benzo(g,h,i)perylene	2.49	0.022	2.42		2.30	0.012	2.26	1.3
1-Methylnaphthalene	2.49	0.22	2.42	0.26	2.30	0.041	2.26	0.17
Benzo[e]pyrene	2.49	0.036	2.42	0.029	2.30	0.031	2.26	1.3
Perylene	2.49		2.42		2.30	0.0089	2.26	0.38
3,6-Dimethylphenanthrene	2.49		2.42		2.30	0	2.26	0.13
1-Methylanthracene	2.49	0.049	2.42	0.040	2.30	0.016	2.26	0.27
1-Methylfluoranthene	2.49		2.42		2.30	0.019	2.26	0.46
1-Methylpyrene	2.49		2.42		2.30	0	2.26	0.13
2,6-Dimethylnaphthalene	2.49	0.16	2.42	0.15	2.30	0.069	2.26	0.21
2,3,5-Trimethylnaphthalene	2.49	0.092	2.42	0.083	2.30	0.044	2.26	0.18
1,1'-Biphenyl	2.49	0.022	2.42		2.30		2.26	
Dibenzofuran	2.49		2.42		2.30		2.26	0.12
1-Methylphenanthrene	2.49	0.084	2.42	0.082	2.30	0.025	2.26	0.14
Dibenzothiophene	2.49	0.029	2.42	0.028	2.30	0.011	2.26	0.13

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations



Table G-1
Summary of Herbicides by Method and Event
Phase I Report Addendum – Additional Data Evaluation

			Numbe	r of Detectio	ons (Parent a	nd Duplicate	· Sample)		Other Meth	or HSM Compared to nods for Total	Compared to WW for	Percent Increase for HSM Compared to LSM for	Percent Increase for HSM Compared to LSM for
	Event/	Total C	oncentration	ns (110/1)	Particula	tes (μg/L)	Dissolve			en Detected by Both	Total Concentrations When Detected by Both	Particulate Concentrations When Detected by Both	Dissolved Concentrations When Detected by Both
Analyte	Attempt	HSM	LSM	WW	HSM	LSM	HSM	LSM	LSM	ww	Methods	Methods	Methods
2,4-DB	All	2	4	2	0	0	2	4	-9%	-46%	-49%		-9%
2,4-D	All	2	2	2	0	0	2	2	-17%	-4%	17%		-17%
2,4,5-T	All	0	1	1	0	0	0	1					
Silvex(2,4,5-TP)	All	2	2	2	0	0	2	2	110%	-47%	-75%		110%
Summary	Market Mark										Alternation and the second and the s		The account
4 Analytes	1/2	0	1.5	0	0	0	0	1.5					
4 Analytes	2/2	0.5	1.0	0	0	0	0.5	1.0	-24%				-24%
4 Analytes	1/3	2.5	2.0	3.5	0	0	2.5	2.0	33%	-32%	-36%		33%
4 Analytes	All	1.0	1.5	1.2	0	0	1.0	1.5	19%	-32%	-36%		19%
Percent of Detected Analytes	All	25%	38%	29%	0%	0%	25%	38%					

Conclusions

The LSM method had the highest number of detections but this result was not statistically significant (see chi-square results).

Where detected in both methods, HSM total concentrations are on average 19% greater than total LSM concentrations.

Where detected in both methods, HSM total concentrations are 32% lower than WW concentrations.

Where detected in both methods, LSM total concentrations are 36% lower than WW concentrations.

Where detected in both methods, HSM dissolved concentrations are on average 19% greater than LSM dissolved concentrations.

No compounds were positively detected using the HSM particulate and LSM particulate analysis.

Abbreviations

μg/L = micrograms per liter
% = percent
HSM = high solids mass
LSM = low solids mass
WW = whole water

Table G-2
Statistical Comparison of the Number of Detected Herbicides by Method and Event
Phase I Report Addendum – Additional Data Evaluation

		of Detectio r Concentr			Fishe	r Exact Test (p-v	alue) ²
Event.	HSM	LSM	ww	Maximum Possible Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW
Event	ПЭМ	LOIVI	VV VV	of Detections		HOW VS. WW	
1/2	0	3	0	8	0.200	1.000	0.200
2/2	1	2	0	8	0.067	1.000	0.467
1/3	5	4	7	8	1.000	0.569	0.282
All	6	9	7	24	0.534	1.000	0.760

Notes

Conclusion

All methods are similar with respect to the number of detected compounds.

Abbreviations

HSM = high solids mass

LSM = low solids mass

WW = whole water

¹ Total number of detections for event includes 2 duplicates and 4 compounds.

² The p-value shown is based on a two-sided Fisher Exact Test. A p-value less than 0.05 is considered evidence of a significant difference among methods compared.

Table G-3
HSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

					HSIV	l Particulate S	Sample Collec	tion				
	E	vent 1 Attempt	2 (7-1-13 PR11	6)	Ev	ent 2 Attempt 2	2 (12-07-13 PR1	35)	Ev	ent 1 Attempt 3	3 (04-30-14 PR1	46)
	PR1CSOC	LYHP-01B	PR1HPC	OUP-01B	PR1CSOC	LYHP-02B	PR1HPC	OUP-02B	PR1CSOC	LYHP-01C	PR1HP	OUP-01C
Wet weight (gram)	50	.39	50	.21	50.23		42	25	49.37		50	.73
% Solids	3	5	32.9		42.5		40.8		48.6		62.3	
Compound Identified	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L
2,4,5-T	0	0	0	0								
2,4-DB												

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

μg/L = micrograms per liter μg/Kg = micrograms per kilogram

Table G-4 HSM Dissolved Analytical Results Phase I Report Addendum – Additional Data Evaluation

					HSI	M Dissolved S	ample Collect	ion						
	Ev	ent 1 Attempt 2	2 (07-01-13 PR11	L7)	Ev	ent 2 Attempt 2	2 (12-07-13 PR13	38)	Eve	ent 1 Attempt 3	(04-30-14 PR14	17)		
	PR1CSOC	LYHD-01B	PR1HDE	OUP-01B	PR1CSOC	LYHD-02B	PR1HDE	OUP-02B	PR1CSOCI	PR1CSOCLYHD-01C PR1HDDUP-01C				
Compound Identified		Sample Result		Sample Result		Sample Result		Sample Result		Sample Result		Sample Result		
Compound Identified	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L		
2,4-DB	0.975		1.015		0.975	0.31	0.975		1.01	0.47	0.960			
2,4-D	0.975		1.015		0.975		0.975		1.01	0.40	0.960	0.41		
2,4,5-T	0.975		1.015		0.975		0.975		1.01	0	0.960	0		
Silvex(2,4,5-TP)	0.975		1.015		0.975		0.975		1.01	0.023	0.960	0.021		

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Table G-5
LSM Particulate Analytical Results
Phase I Report Addendum – Additional Data Evaluation

					LSM	Particulate S	ample Colle	ction				
	Eve	nt 1 Attempt 2	. (07-05-13 PR:	119)	Eve	nt 2 Attempt 2	(12-11-13 PR	140)	Even	t 1 Attempt 3	(05-05-2015 PI	R149)
	PR1CSOC	CLYLP-01B	PR1LPD	OUP-01B	PR1CSOC	CLYLP-02B	PR1LPD	UP-02B	PR1CSOC	CLYLP-01C	PR1LPDUP-01C	
TSS (mg/L)	64	1.8	67.1		8	.4	13	3.5	8	3	:	8
Total Liters Filtered (L)	0.9	984	0.994		1.0)42	1.0)10	1.0)53	1.0	027
Compound Identified	Weight gram	Sample Result ug/Kg	Weight gram	Sample Result ug/Kg	Weight gram	Sample Result ug/Kg	Weight gram	Sample Result ug/Kg	Weight gram	Sample Result ug/Kg	Weight gram	Sample Result ug/Kg
N/A	0.0640		0.0666		0.0088		0.0136		0.00842		0.00822	

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

μg/Kg = micrograms per kilogram mg/L = milligrams per liter

Table G-6
LSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

					LSIV	Dissolved Sa	ample Collect	ion				
	Eve	ent 1 Attempt 2	2 (07-05-13 PR1	20)	Eve	ent 2 Attempt 2	2 (12-11-13 PR1	41)	Eve	ent 1 Attempt 3	3 (05-05-14 PR1	50)
	PR1CSOC	LYLD-01B	PR1LDD	UP-01B	PR1CSOCLYLD-02B		PR1LDDUP-02B		PR1CSOCLYLD-01C		PR1LDDUP-01C	
		Sample Result		Sample Result		Sample Result		Sample Result		Sample Result		Sample Result
Compound Identified	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L
2,4-DB	1.012	0.45	0.994	1	1.04		1.01	0.41	1.00		1.035	0.44
2,4-D	1.012		0.994		1.04		1.01		1.00	0.47	1.035	0.51
2,4,5-T	1.012		0.994		1.04		1.01	0.21	1.00	0	1.035	0
Silvex(2,4,5-TP)	1.012	0.02	0.994		1.04		1.01		1.00		1.035	0.021

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>

Table G-7
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

					Wh	ole Water Sa	mple Collecti	on					
	Eve	ent 1 Attepmt 2	2 (07-01-13 PR1	15)	Eve	ent 2 Attempt 2	(12-07-13 PR1	34)	Eve	ent 1 Attempt 3	(04-30-14 PR1	80-14 PR145)	
	PR1CSOCL	YWW-01B	PR1WW	DUP-01B	PR1CSOCLYWW-02B		PR1WW	DUP-02B	PR1CSOCL	YWW-02B	PR1WWDUP-02B		
Compound Identified	Volume Liters	Sample Result Volume Liters ug/L Vo		Sample Result lume Liters ug/L		Sample Result ug/L	Volume Liters	Sample Result ug/L	Sample Result Volume Liters ug/L		Volume Liters	Sample Result ug/L	
2,4-DB	0.990		0.980		1.025		1.015		0.935	0.59	1.0	0.28	
2,4-D	0.990		0.980		1.025		1.015		0.935	0.36	1.0	0.48	
2,4,5-T	0.990		0.980		1.025		1.015		0.935	0	1.0	0.1	
Silvex(2,4,5-TP)	0.990		0.980		1.025		1.015		0.935	0.051	1.0	0.032	

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Appendix H

Data Evaluation Summaries and Analytical Results - Cyanide

Table H-1
Summary of Cyanide by Method and Event
Phase I Report Addendum – Additional Data Evaluation

		Nui	mber of [etections	(Parent a	and Dupli	icate Sam	ple)	Compared to	rease for HSM Other Methods	Percent Increase for LSM Compared to WW for Total	l	Percent Increase for HSM Compared to LSM for
	Event/	Total	Concentr (μg/L)	ations	Partice (µg		Dissolve	ed (µg/L)	for Total Cond Detected by	entrations When Both Methods	WW for Total Concentrations When Detected by Both	Concentrations When Detected by Both	Dissolved Concentrations When Detected by Both
Analyte	Attempt	нѕм	LSM	ww	нѕм	LSM	нѕм	LSM	LSM	ww	Methods	Methods	Methods
CN	All	4	NA	4	4	NA	2	NA					
Summary													
1 Analyte	1/2	1.0	NA	1.0	1.0	NA	1.0	NA		12%			
1 Analyte	2/2	1.0	NA	1.0	1.0	NA	0.0	NA		-98%			
1 Analyte	Ali	1.0	NA	1.0	1.0	NA	0.5	NA		-43%			
Percent of Detected Analytes	Ali	100%	NA	100%	100%	NA	50%	NA					

Conclusions

The frequency of detection was same (equal) for HSM total and whole water concentrations.

Where detected in both methods, HSM total concentrations are 43% lower than WW concentrations. However, it should be noted that total concentrations in Event 1/2 were similar between HSM total and WW but WW concentrations were of a magnitue approximately 10 times greater than HSM in Event 2/2.

<u>Abbreviations</u>

 μ g/L = micrograms per liter

% = percent

HSM = high solids mass

LSM = low solids mass

WW = whole water

CN = cyanide

NA = Cyanide was not analyzed for LSM particulate/dissolved samples.

Table H-2 **HSM Particulate Analytical Results** Phase I Report Addendum - Additional Data Evaluation

			HSI	M Particulate S	Sample Collect	ion		
		Event 1 Attempt	2 (7-1-13 PR116		E,	vent 2 Attempt 2	! (12-07-13 PR13!	5)
	PR1CSOC	LYHP-01B	PR1HPE	UP-01B	PR1CSOC	LYHP-02B	PR1HPC	UP-02B
Wet weight (gram)	2.	03	3 2.		1.01		í	L
% Solids	26	5.7	26	5.3	42	2.5	40.8	
Compound Identified	Sample Result mg/Kg	Converted Sample Result ug/L	Sample Result mg/Kg	Converted Sample Result ug/L	Sample Result mg/Kg	Converted Sample Result ug/L	Sample Result mg/Kg	Converted Sample Result ug/L
Cyanide	5.8	0.138	6.400	0.150	2.4	0.0905	1.60	0.0579

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

μg/L = micrograms per liter

mg/KG = milligrams per kilogram

Table H-3
HSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

		HSM Dissolved Sample Collection									
	E	vent 1 Attempt 2	(07-01-13 PR11	7)	E	Event 2 Attempt 2 (12-07-13 PR138)					
	PR1CSOC	LYHD-01B	PR1HDI	OUP-01B	PR1CSOC	LYHD-02B	PR1HDI	OUP-02B			
Compound Identified	Volume Liters			Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L			
Cyanide	0.05	31.3	0.05	31.6	0.05		0.05				

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Table H-4
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

		Whole Water Sample Collection									
	l	Event 1 Attepmt 2	(07-01-13 PR115		Event 2 Attempt 2 (12-07-134 PR134)						
1	PR1CSOCL	YWW-01B	PR1WW	DUP-01B	PR1CSOCL	YWW-01B	PR1WWDUP-01B				
Compound Identified	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result			
Compound identified	Liters	ug/L	Liters	ug/L	Liters	ug/L	Liters	ug/L			
Cyanide	0.05	29.3	29.3 0.05 27.2		0.05	3.8	0.05	2.3			

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Α	ppe	nd	ix	I
				-

Data Evaluation Summaries and Analytical Results – VOCs

Table I-1
Summary of VOCs by Method and Event
Phase I Report Addendum – Additional Data Evaluation

			Number of Detections (Parent and Duplicate Sample)						Percent Increa Compared Methods f Concentratio	to Other or Total ons When	Percent Increase for LSM Compared to WW for Total	Percent Increase for HSM Compared to LSM for Particulate	to LSM for
	l _								Detected I	•	Concentrations	Concentrations	Concentrations
	Event/		ncentratio		Particula		Dissolved	··· ·	Metho		When Detected by	When Detected by	When Detected by
Analyte	Attempt	HSM	LSM	ww	HSM	LSM	HSM	LSM	LSM	ww	Both Methods	Both Methods	Both Methods
1,4-Dichlorobenzene	All	2	NA	0	2	NA	0	NA					
Summary													
6 Analytes	1/2	1.0	NA	0.0	1.0	NA	0.0	NA					
6 Analytes	2/1	0.0	NA	0.0	0.0	NA	0.0	NA					
6 Analytes	All	0.5	NA	0.0	0.5	NA	0.0	NA					
Percent of Detected Analytes	All	8%	NA	0%	8%	NA	0%	NA					

Conclusions

Samples were analyzed for a total of 6 VOC compounds, however, only VOC compounds that were positively identified during analysis are presented. Positive results were reported for HSM particulate analysis only.

Abbreviations

μg/L = micrograms per liter

% = percent

HSM = high solids mass

LSM = low solids mass

WW = whole water

NA = Volatile organic compounds (VOCs) were not analyzed for LSM particulate/dissolved samples.

Table I-2
Statistical Comparison of the Number of Detected VOCs by Method and Event
Phase I Report Addendum – Additional Data Evaluation

	Number of Detections (Total Water Concentration)				Fishe	alue) ²	
				Maximum Possible			
Event	HSM	LSM	ww	Number of Detections ¹	HSM vs. LSM	HSM vs. WW	LSM vs. WW
1/2	2	NA	0	12	NA	0.478	NA
2/2	0	NA	0	12	NA	1.000	NA
All	2	NA	0	24	NA	0.489	NA

Notes

Conclusion

HSM had only 2 detections while WW had none.

LSM was not evaluated.

Differences were not statistically significant.

Abbreviations

HSM = high solids mass

LSM = low solids mass

WW = whole water

NA = Not analyzed.

¹ Total number of detections for event includes 2 duplicates and 6 compounds.

² The p-value shown is based on a two-sided Fisher Exact Test. A p-value less than 0.05 is considered evidence of a significant difference among methods compared.

Table I-3 HSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

		HSM Particulate Sample Collection									
		Event 1 Attempt	2 (7-1-13 PR116)			Event 2 Attempt 1 (10-07-13 PR129)					
	PR1CSOC	CSOCLYHP-01B PR1HPDUP-01B			PR1CSOCI	YHP-02A2	PR1HPDUP-02A2				
Wet weight (gram)	2.	24	3.26		2.23		1.94				
% Solids	2	27	26		34		34				
Compound Identified	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L	Sample Result ug/Kg	Converted Sample Result ug/L			
1,4-Dichlorobenzene	47	0.00113	15	0.000347							
Chlorobenzene	0	0	0	0							

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

 μ g/L = micrograms per liter

μg/KG = micrograms per kilogram

Table I-4
HSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

		HSM Dissolved Sample Collection								
	E	Event 1 Attempt 2 (07-01-13 PR117) Event 2 Attempt 1 (10-								
	PR1CSO	CLYHD-01B	PR1H	IDDUP-01B	PR1CSC	OCLYHD-02A	PR1H	DDUP-02A		
Compound Identified	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L	Volume Liters	Sample Result ug/L		
1,4-Dichlorobenzene	0.025	0	0.025	0	0.025	0	0.025	0		

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Table I-5
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

ſ				W	hole Water Sa	ımple Collecti	on				
ı		Εν	ent 1 Attepmt 2	(07-01-13 PR1:	15)	Event 2 Attempt 1 (10-07-13 PR128)					
ı		PR1CSOCI	LYWW-01B	PR1WW	DUP-01B	PR1CSOCI	.YWW-02A	PR1WWDUP-02A			
ı											
ı	Compound Identified		Sample Result		Sample Result		Sample Result		Sample Result		
L		Volume Liters ug/L		Volume Liters	ug/L	Volume Liters	ug/L	Volume Liters	ug/L		
	1,4-Dichlorobenzene	0.025	0	0.025	0	0.025	0	0.025	0.0000		

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

Abbreviations

Δr	pen	dix	ı.
7	pen	UIA	J

Data Evaluation Summaries and Analytical Results – TEPH

Table J-1
Summary of Total EPH by Method and Event
Phase I Report Addendum – Additional Data Evaluation

		Number of Detections (Parent and Duplicate Sample)						÷)	Percent Incre			Percent Increase for HSM Compared to	Percent Increase for HSM Compared to
									for Total Cor	for Total Concentrations		LSM for Particulate	LSM for Dissolved
									When Detec	ted by Both	Concentrations	Concentrations	Concentrations When
	Event/	Total Con	centratio	ns (mg/L)	Particulat	es (mg/L)	Dissolve	d (mg/L)	Meti	hods	When Detected by	When Detected by	Detected by Both
Analyte	Attempt	HSM	LSM	ww	HSM	LSM	HSM	LSM	LSM	ww	Both Methods	Both Methods	Methods
TEPH	All	4	NA	4	4	NA	2	NA					
Summary													
1 Analyte	1/2	1.0	NA	1.0	1.0	NA	1.0	NA		-22%			
1 Analyte	2/2	1.0	NA	1.0	1.0	NA	0.0	NA		-88%			
1 Analyte	Ali	1.0	NA	1.0	1.0	NA	0.5	NA		-55%			
Percent of Detected Analytes	Ali	100%	NA	100%	100%	NA	50%	NA					

Conclusions

The frequency of detection was same (equal) for HSM total and whole water concentrations.

Where detected in both methods, HSM total concentrations are 55% lower than WW concentrations.

NA = total extrctable petroleum hydrocarbon (TEPH) was not analyzed for LSM particulate/dissolved samples.

<u>Abbreviations</u>

mg/L = milligrams per liter

% = percent

HSM = high solids mass

LSM = low solids mass

WW = whole water

Table J-2 HSM Particulate Analytical Results Phase I Report Addendum – Additional Data Evaluation

		HSM Particulate Sample Collection										
		Event 1 Attempt	2 (7-1-13 PR116)		Event 2 Attempt 2 (12-07-13 PR135)							
	PR1CSOC	LYHP-01B	PR1HPE	OUP-01B	PR1CSOC	CLYHP-02B	PR1HPDUP-02B					
Wet weight (gram)	30	.24	30.18		30.03		29	.43				
% Solids	3	15	32.9		42.5		40	0.8				
Compound Identified	Sample Result mg/Kg	Converted Sample Result mg/L	Sample Result mg/Kg	Converted Sample Result mg/L	Sample Result mg/Kg	Converted Sample Result mg/L	Sample Result mg/Kg	Converted Sample Result mg/L				
ТЕРН	13,000	0.405	13,000	0.380	13,000	0.491	7,700	0.279				

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>

mg/L = milligrams per liter

mg/Kg = milligrams per kilogram

TEPH = total extrctable petroleum hydrocarbon

Table J-3
HSM Dissolved Analytical Results
Phase I Report Addendum – Additional Data Evaluation

	HSM Dissolved Sample Collection Event 1 Attempt 2 (07-01-13 PR117) Event 2 Attempt 2 (12-07-13 PR138)									
	PR1CSOC	LYHD-01B	PR1HDI	OUP-01B	PR1CSOCLYHD-02B		PR1HDDUP-02B			
Compound Identified	Volume Sample Result Liters mg/L		Volume Liters	Sample Result mg/L	Volume Liters	Sample Result mg/L	Volume Liters	Sample Result mg/L		
TEPH	0.995	5.6	1.045	3.5	1.055		1.030			

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a ND result.

Abbreviations

mg/L = milligrams per liter

TEPH = total extrctable petroleum hydrocarbon

Table J-4
Whole Water Analytical Results
Phase I Report Addendum – Additional Data Evaluation

Compound Identified	Whole Water Sample Collection							
	Event 1 Attepmt 2 (07-01-13 PR115)				Event 2 Attempt 2 (12-07-13 PR134)			
	PR1CSOCLYWW-01B		PR1WWDUP-01B		PR1CSOCLYWW-02B		PR1WWDUP-02B	
	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result	Volume	Sample Result
	Liters	mg/L	Liters	mg/L	Liters	mg/L	Liters	mg/L
TEPH	1.020	5.0	1.060	7.7	1.050	2.22	0.985	4.200

A "0" value in the sample result column represents a result that was qualified by the lab as "G". A "G" qualifier indicates the presence of a compound that meets the identification criteria; the result is below the PQL but above the method detection limit (MDL) or estimated detection limit (EDL), where appropriate.

A "null" value in the sample result column represents a non-detect (ND) result.

<u>Abbreviations</u>

mg/L = milligrams per liter

TEPH = total extrctable petroleum hydrocarbon